

Prepared in cooperation with the METRO WASTEWATER RECLAMATION DISTRICT

BIOSOLIDS, SOIL, CROP, GROUND-WATER, AND STREAMBED-SEDIMENT DATA FOR A BIOSOLIDS-APPLICATION AREA NEAR DEER TRAIL, COLORADO, 2000

Open-File Report 03-400



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Application Area Near Deer Trail, Colorado, 2000
By Tracy J.B. Yager, David B. Smith, James G. Crock, and Michael R. Stevens

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U.S. Department of the Interior

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Conversion Factors

Multiply	by	To obtain
acre	0.4047	hectares (ha.)
centimeter (cm)	0.3937	inch
foot (ft)	0.3048	meter
gallon (gal)	3.785	liter
gram (g)	0.035	ounce
inch	2.54	centimeter
liter (L)	0.2642	gallon (gal)
micrometer (μm)	0.00003937	inch
mile (mi)	1.609	kilometer
milliliter (mL)	0.0610	cubic inch
millimeter (mm)	0.03937	inch
square mile (mi ²)	2.590	square kilometer

Temperature in degrees Celsius (°C) can be converted to degrees Fahrenheit (°F) using the formula $^{\circ}F = (1.8 \text{ x}^{\circ}\text{C}) + 32.$

Temperature in degrees Fahrenheit (°F) may be converted to degrees Celsius (°C) using the formula $^{\circ}C = (^{\circ}F - 32)/1.8$

Vertical coordinate information is referenced to National Geodetic Vertical Datum of 1929 (NGVD 29); horizontal coordinate information is referenced to North American Datum of 1927 (NAD 27) except as noted.

ADDITIONAL ABBREVIATIONS

gal/min	gallon per minute
mg	milligrams
mg/g	milligrams per gram
μg/L	micrograms per liter
μS/cm	microsiemens per centimeter at 25 degrees Celsius
mg/kg	milligrams per kilogram
mg/L	milligrams per liter
ρ C i/L	picocuries per liter
ρ C i/g	picocuries per gram
ROE	residue on evaporation

Biosolids, Soil, Crop, Ground-Water, and Streambed-Sediment Data for a Biosolids-Application Area Near Deer Trail, Colorado, 2000

By Tracy J.B. Yager, David B. Smith, James G. Crock, and Michael R. Stevens

Abstract

In January 1999, the U.S. Geological Survey began an expanded monitoring program near Deer Trail, Colorado, in cooperation with the Metro Wastewater Reclamation District and the North Kiowa Bijou Groundwater Management District. Monitoring components were biosolids, soils, crops, ground water, and streambed sediments. The monitoring program addresses concerns from the public about chemical effects from applications of biosolids to agricultural land in the Deer Trail area. Constituents of primary concern to the public are arsenic, cadmium, copper, lead, mercury, molybdenum, nickel, selenium, zinc, plutonium, and gross alpha and beta activity, and they are included for all monitoring components. This report presents chemical data from the second year of the monitoring program, January-December 2000, for biosolids, crops, alluvial and bedrock ground water, and streambed sediments. The ground-water section also includes climate data, water levels, and results of statistical testing of selected data for trends and for exceedance of Colorado regulatory standards. The chemical data included are for the constituents of highest concern to the public as well as for many other constituents.

Introduction

Since 1993, the Metro Wastewater Reclamation District (MWRD) has been applying biosolids resulting from municipal sewage treatment in Denver, Colo., to their property near Deer Trail, Colo. The biosolids are transported about 75 mi east from Denver to the MWRD property and are applied to nonirrigated farmland. From 1993 to 1999, the U.S. Geological Survey (USGS), in cooperation with the MWRD, monitored the quality of shallow ground water on the MWRD central property (fig. 1 in the Data Section at the back of the report), which encompassed about 15 mi² and was the first property the MWRD purchased near Deer Trail. In 1995, the

MWRD traded some of the property and acquired additional property in the same area. The new property consisted of about 14.5 mi² known as the north property and about 50 mi² known as the south property. In 1999, the three MWRD properties together, known as the METROGRO Farm, encompassed almost 70 mi² of farmland, including land in Arapahoe and Elbert Counties. The three MWRD properties and surrounding private property are hereinafter referred to as the study area (fig. 1).

The study area is on the eastern plains of Colorado about 10 mi east of Deer Trail. The study area is on the eastern margin of the Denver Basin, a bowl-shaped sequence of sedimentary rocks that was formed in an ocean or near-ocean environment. The geology of the study area consists of interbedded shale, siltstone, and sandstone, which may be overlain by clay, windblown silt and sand, or alluvial sand and gravel (Sharps, 1980; Major and others, 1983; Robson and Banta, 1995). The primary water-supply aquifer is the Laramie-Fox Hills aquifer, which is a bedrock aquifer that ranges from 0 to about 200 ft thick in the study area and is the bottom aguifer in the Denver Basin aguifer sequence (Robson and others, 1981; Robson and Banta, 1995). Multiple alluvial aquifers are present in the study area. These aguifers are associated with the surficial drainage network but contain water of variable quality, are of limited extent, and generally yield little water (Stevens and others, 2003; Yager and Arnold, 2003). The study area is within the South Platte River drainage basin; all streams in this area drain northward to the South Platte River (U.S. Geological Survey, 1974; Seaber and others, 1987; Yager and Arnold, 2003). Short segments of some of the streams are intermittent, but in general, the streams are ephemeral and flow only after storms. No surface water flows off the MWRD properties except after storms. Most ponds in the area have been created by detention structures. Soils in the area generally are sandy or loamy on flood plains and stream terraces, clayey to loamy on gently sloping to rolling uplands, and sandy and shaley on steeper uplands (Larsen and others, 1966; Larsen and Brown, 1971).

Land use in the study area was historically rangeland or cropland and pasture (U.S. Geological Survey, 1980). Some petroleum exploration was done in the area (Drew and others, 1979), but no oil or gas production took place within the study area during 2000. Land use during 2000 was rangeland or cropland. Cattle and sheep are the primary domesticated animals grazing the area, and wheat is the primary crop. Farmland is not irrigated. Land use on the MWRD properties during 2000 was primarily cropland (with biosolids applied as a fertilizer) and some rangeland.

Public concern about applications of biosolids to farmland increased after the MWRD agreed to accept treated ground water from the Lowry Landfill Superfund site in Denver. The concern was that water from the Lowry Superfund site might contain radionuclides that would then contaminate the MWRD biosolids. In January 1999, the USGS began a new monitoring program in cooperation with the MWRD and the North Kiowa Bijou Groundwater Management District. The USGS refers to the new monitoring program (1999–2005) as the "expanded monitoring program."

The expanded monitoring program near Deer Trail is distinct from, but builds on, the previous monitoring program in which the USGS monitored the quality of shallow ground water on the MWRD central property (1993–99). Relative to the previous program, the expanded program includes a larger study area (fig. 1) (all three MWRD properties and private-property locations), more monitoring components (biosolids, soils, crops, and streambed sediments in addition to ground water), a more comprehensive list of chemical constituents, expanded statistical analyses of data, and an extended monitoring period (1999–2005). Both programs use USGS and MWRD funds. Both programs are designed, accomplished, and interpreted independently by the USGS, and quality-assured USGS data and reports are released to the public and the MWRD at the same time.

Biosolids are applied by the MWRD to their properties near Deer Trail according to agronomic loading rates. Landapplied biosolids must meet Colorado biosolids regulations for metals and radioactivity; otherwise, soils could become overloaded. Soil quality either can be improved by biosolids applications through increased nutrients and organic matter or degraded through accumulation of excessive nutrients or metals. Pesticides, herbicides, and other fertilizers also may have been applied to the MWRD properties in the past, but less information is available about these applications.

Animal waste related to grazing domestic livestock and applications of pesticides, herbicides, and fertilizers (including biosolids) can affect soil quality, crops, water quality in alluvial and bedrock aquifers, and streambed-sediment chemistry. Water quality can be affected directly by contaminated recharge water or by infiltration of water through contaminated soils or sediments (remobilization). Water quality can be affected indirectly by tillling that mobilizes or mixes subsurface chemical constituents or by contributions to natural processes such as nitrification. Contaminated ground water or surface water could contaminate other aquifers (such as

bedrock water-supply aquifers or alluvial aquifers), other surface-water bodies (ponds or streams), or streambed sediment.

The expanded monitoring program near Deer Trail addresses these concerns about biosolids applications and other farming-related effects on the environment and should increase scientific insight about Denver Basin hydrology. The objectives of this USGS program are to: (1) evaluate the combined effects of biosolids applications, land use, and natural processes on soils, crops, the bedrock aquifer, alluvial aquifers, and streambed sediments by comparing chemical data to (a) regulatory standards, (b) data from a site where biosolids are not applied (a control site), or (c) earlier data from the same site (trends); (2) monitor biosolids for trace elements and radioactivity and compare trace-element concentrations and radioactivity with regulatory standards; and (3) characterize the hydrology of the study area. The monitoring of each component (such as soil or ground water) is a stand-alone study that includes radioactivity analyses because of public concerns about effects from the Lowry Landfill Superfund site. More detailed information about the monitoring of each component is included later in this report.

Purpose and Scope

The purpose of this report is to present information from the expanded monitoring program near Deer Trail for 2000 (January through December). This report presents data for all monitoring components of the program except soils. The report includes information about biosolids, soils, crops, ground water (alluvial and bedrock), and streambed sediment. The ground-water sections include climate data, lithologic descriptions and well-completion diagrams, hydrologic data (depth to ground water), water-quality data (chemistry and field measurements), and results of statistical testing of selected data for exceedance of regulatory standards and trends. This report does not include the hydrogeologic structure maps that were done as part of the bedrock ground-water monitoring component of the program. The structure maps were used to select bedrock-aquifer monitoring locations for the expanded monitoring program. The structure maps, along with a more detailed discussion of the hydrogeology of the region, are included in an interpretive USGS report by Yager and Arnold (2003).

This report is organized by monitoring component because each component (such as soil or ground water) is monitored as a separate study. For each monitoring component, the specific objectives, scope, approach, analytical results, quality-assurance information, and a discussion are included. Data in this report for January–June 2000 were collected by the USGS before any water transfer from the Lowry Landfill Superfund site to the MWRD treatment plant. The water transfer from the Lowry Landfill Superfund site to the MWRD began in July 2000. Therefore, some of the data in this report, along with all the data in the 1999 report (Stevens and others, 2003), provide a geochemical baseline for

each monitoring component prior to the Lowry water transfer. These baselines will enable the USGS to recognize and quantify potential chemical changes in each monitoring component from the Lowry water transfer.

Acknowledgments

The USGS thanks all private landowners for allowing access to their properties for data collection. The USGS especially thanks the Price and Weisensee families and the MWRD for allowing USGS instrument or well installations on their property and the Kalcevic family for timely sediment information after storms and for allowing streambed-sediment sampling on their property.

Biosolids

Biosolids are solid organic matter recovered from a sewage-treatment process that meet State and Federal regulatory criteria for beneficial use, such as for fertilizer. Land-applied biosolids must meet or exceed Grade II, Class B criteria (Colorado Department of Public Health and Environment, 1998). Grade I exceeds Grade II. The MWRD applies Grade I, Class B biosolids to its properties near Deer Trail. The biosolids-application areas, dates of application, and application rates provided by the MWRD for its properties near Deer Trail are listed in table 1 (located in the Data Section at the back of the report); application areas (called "Destination Codes") are marked DC and shown in figure 2 (in the Data Section at the back of the report).

Objectives of Monitoring Biosolids

The biosolids must meet regulatory standards for trace elements and radioactivity. Exceeding these standards could adversely affect the quality of soil on which the biosolids are applied and could alter MWRD plans for the application of biosolids in Arapahoe and Elbert Counties. The composition of the biosolids was monitored to provide an independently determined data set against which the MWRD chemical analyses and the regulatory standards for biosolids can be compared. The data also will constitute a chemical baseline against which any future change in the concentration of constituents analyzed for in this study may be recognized, measured, and compared.

Approach for Monitoring Biosolids

In 2000, the USGS continued the protocol established in 1999 for monitoring MWRD biosolids for concentrations of arsenic, cadmium, copper, lead, mercury, molybdenum, nickel, selenium, zinc, plutonium, and gross alpha and beta activity.

Radioactivity analyses were included in response to public concerns that biosolids radioactivity could increase from the transfer of water from the Lowry Landfill Superfund site.

Biosolids samples were collected directly from the MWRD facility in Denver rather than from individual trucks or fields near Deer Trail to enable the USGS to obtain a more representative sample. Normal annual sampling protocol calls for one biosolids sample to be collected each quarter. In late July 2000, however, MWRD began receiving treated water from the Lowry Landfill Superfund site. Therefore, in August 2000, the USGS initiated monthly sampling for the remainder of the year. In 2000, samples were collected for March, June, August, September, October, November, and December. The samples were prepared and analyzed at the chemical laboratories of the USGS Mineral Resources Program in Denver. The concentrations in the samples were compared to applicable Colorado standards for biosolids (Colorado Department of Public Health and Environment, 1998).

Sampling Methods for Biosolids

Each biosolids sample is a 24-hour composite of 12 subsamples collected about every 2 hours by MWRD personnel at the MWRD facility. The subsamples were collected from the conveyor belt that transfers the biosolids into the transport trucks. Each sample was delivered to the USGS in two acidwashed, rinsed, 1-gallon plastic bottles.

Analytical Methods for Biosolids

The biosolids material was air dried and then ground to less than 150 μm before chemical analysis. The biosolids samples were processed and analyzed for trace elements at the laboratories of the USGS Mineral Resources Program in Denver and analyzed for radioactivity at a commercial laboratory, Acculabs in Colorado. The methods used to analyze the biosolids for each constituent are listed in table 2 (located in the Data Section at the back of the report).

Quality Assurance for Biosolids

The purpose of the quality-assurance program developed for the biosolids monitoring component was to ensure the analytical results were within acceptable limits of both precision (the reproducibility of results) and accuracy (the degree of conformity of results for a sample having known concentrations). The precision was determined by analyzing the same biosolids sample multiple times, and accuracy was determined by analyzing National Institute of Standards and Technology (NIST) standard reference material SRM 2781, a domestic sludge. This standard reference material (SRM) was prepared by the NIST from material collected at the MWRD treatment plant in Denver. SRM 2781 has been analyzed extensively by many laboratories throughout the world, and the NIST has certified an acceptable range of values for various constituents

in the SRM. The constituents include those of interest in this study. Each biosolids sample was submitted to the laboratories with a sample of the SRM. If the analytical results for the constituent of interest in the SRM were within the acceptable range, the results for the biosolids samples were accepted.

In 2002, the USGS became concerned about the gross alpha data for the biosolids samples. For the March 1999 through June 2000 samples, the gross alpha data from Acculabs for the NIST standard reference material 2781 (domestic sludge) ranged from 27-37 pCi/g. For the August 2000 through August 2001 samples, the gross alpha data ranged from 37- 60 pCi/g. This shift to higher values for the same standard-reference material indicates possible increasing analytical bias that could be present in the gross alpha data for the biosolids samples collected from the MWRD. Additional information about these analyses is not available because Acculabs went out of business in early 2002. Therefore, the USGS submitted split samples from a subset of the biosolids samples and NIST standard reference material to a different laboratory in an attempt to reconcile this issue. A split of the June 2000 biosolids sample and three splits of the NIST standard reference material 2781 were analyzed for radioactivity in 2002 by Severn Trent Laboratory (formerly Quanterra Analytical Services) in Richland, Wash., under a contract with the USGS National Water Quality Laboratory (NWQL).

Biosolids Data

Summaries of all the chemical analyses for trace-element concentrations and radioactivity data (gross alpha activity, gross beta activity, and plutonium concentration) for the biosolids samples collected in 2000 are listed in table 3 (in the Data Section at the back of the report). The tables also list the maximum allowable concentrations for Grade I biosolids. Radioactivity data for the same samples from two different laboratories are listed in table 4 (at the back of the report).

Discussion of Biosolids Data

All trace-element concentrations were less than the maximum allowable concentrations established for Grade I biosolids. The data from Severn Trent Laboratory (STL) compare with the Acculabs data from 2000. No significant analytical bias or variability likely is present in the 2000 biosolids data from Acculabs.

Soils

Biosolids can contain elevated concentrations of certain trace constituents. Therefore, the application of biosolids to farmland has caused public concern regarding the potential short-term and long-term effects on soil quality.

Objectives of Monitoring Soils

Soils were monitored for trace elements and plutonium and gross alpha and gross beta activity to establish independent geochemical data sets for the composition of soil before and after the application of biosolids. The data will enable the USGS to recognize and quantify significant changes in soil composition caused by the application of biosolids to agricultural soils or by other natural or human-induced processes.

Approach for Monitoring Soils

In August 1999, the USGS began monitoring soils on two sites, one site on MWRD property in Arapahoe County and one site on MWRD property in Elbert County. The sites were monitored for arsenic, cadmium, copper, lead, mercury, molybdenum, nickel, selenium, zinc, plutonium, and gross alpha and beta activity. Radioactivity analyses were included in response to public concerns that biosolids radioactivity could increase from the transfer of water from the Lowry Landfill Superfund site. Soil samples were collected once during 1999, before the application of biosolids to monitoring sites, and the data were reported in quarterly reports and in the annual data report for 1999 (Stevens and others, 2003). Soil monitoring will continue through two cycles of biosolids application and crop harvest, and soil sampling will be done shortly after each harvest.

Site Selection for Monitoring Soils

Sites were selected on MWRD properties where biosolids have never been applied. One site was selected on the MWRD north property in Arapahoe County, and one site was selected on the MWRD south property in Elbert County. The Araphoe County site is located in T. 4 S., R. 58 W., sec. 22 and lies about 0.25 mi west of Badger Creek (fig. 3). The Elbert County site is located in T. 6 S., R. 57 W., sec. 8 and lies immediately west of Beaver Creek (fig. 4).

Sampling Methods for Soils

The sampling protocol was designed to answer the following question: What is the average composition of the top 12 inches of soil in each of the six 20-acre fields? To address this question, a standard soil auger was used to collect samples to a depth of 12 inches according to a systematic grid pattern. For each of the two fields to which biosolids will be applied, 36 subsamples are collected on approximately 133-ft centers. A similar grid is used to collect 36 subsamples from the southern "control" field on the Arapahoe County site. For the remaining three "control" fields, 30 subsamples are collected at approximately 155-ft centers.

Analytical Methods for Soils

All soil subsamples are air dried at ambient temperature in the laboratories of the USGS Mineral Resources Program in Denver. Each of the dried subsamples is disaggregated and sieved to less than 2 mm. This minus-2-mm material then is ground to less than 150 μ m. Splits of each subsample are taken for archival storage, and the subsamples for each field are composited into one sample for chemical analysis. The composite soil samples, each representing one 20-acre field, are analyzed by the methods listed in table 2.

Quality Assurance for Soils

The accuracy of the soil analysis was ensured by the analysis of NIST SRM 2709, an agricultural soil. Separate splits of this SRM were randomly placed among the soil samples and submitted to the laboratories. If the analytical results for the constituent of interest in the SRM are within an acceptable range, analytical results for the soil samples are accepted. Separate splits of each composited soil sample are analyzed independently and the results averaged to determine the concentration reported for a given constituent.

Soil Data

In 2000, the Elbert County monitoring site was sampled in late November after the wheat harvest. The Arapahoe County site was not sampled until June 2001. The samples from both sites were submitted to the laboratories for analysis in September 2001. Data for both these sites will be presented in progress reports as the data become available and in the annual project data report for 2001.

Crops

As previously mentioned, biosolids can contain elevated concentrations of certain trace elements. The application of biosolids to farmland on which grain crops are grown that will eventually be consumed by animals or humans has led to public concern about the composition of the crops grown on the fields receiving biosolids. Both the Arapahoe and Elbert County sites were originally planted in wheat during the winter of 1999–2000. The wheat on the Elbert County site reached maturity and was harvested in July 2000. The wheat crop on the Arapahoe County site failed and the field was plowed under in May 2000. Approximately the western one-half of the site was replanted with millet during spring 2000; the millet reached maturity and was harvested in September 2000.

Objectives of Monitoring Crops

Crops were monitored for trace elements (and selected samples were monitored for plutonium and gross alpha and gross beta activity) in order to establish independent chemical data sets for the composition of the crops before and after the application of biosolids. The data will enable the USGS to recognize and quantify significant changes in crop composition caused by the application of biosolids to agricultural soils or by other natural or human-induced processes.

Approach for Monitoring Crops

In the summer of 2000, the USGS began monitoring crops grown on the same two sites where soils were monitored. One of these sites is on MWRD property in Arapahoe County, and one site is on MWRD property in Elbert County. The crops were monitored for arsenic, cadmium, copper, lead, mercury, molybdenum, nickel, selenium, and zinc. One sample from the Elbert County site was analyzed for plutonium and gross alpha and beta activity in response to public concerns that biosolids radioactivity could increase because of the transfer of water from the Lowry Landfill Superfund site.

Crops grown on fields that receive biosolids applications were monitored along with crops grown on fields that do not receive biosolids applications. The crops from fields that do not receive biosolids applications were used as a reference for comparison. Each of the two crop- and soil-monitoring sites consisted of three 20-acre (933-ft by 933-ft) fields separated by 100-ft buffer zones (figs. 3 and 4, in the Data Section at the back of the report). In 1999, the center 20-acre field at each site received a single biosolids application after the initial soil sampling. The other two 20-acre fields at each site will not receive biosolids and will be used as "control" fields to determine the natural variability of soil and crop composition for the duration of the study. All three 20-acre fields at each site are farmed the same way as the rest of the MWRD property and have crops planted and harvested. Crop samples from each of the six fields were collected in the summer of 2000. Data will be compared after each sampling and at the conclusion of the study to determine how the concentrations of the constituents of interest vary with time.

Site Selection for Monitoring Crops

The soil-monitoring fields (figs. 3 and 4) also will be used for monitoring crops. Crops are planted on each of the six 20-acre soil-monitoring fields. Sites were selected on MWRD properties where biosolids have never been applied. One site was selected on the MWRD north property in Arapahoe County, and one site was selected on the MWRD south property in Elbert County. The Arapahoe County site is located in T. 4 S., R. 58 W., sec. 22 and lies about 0.25 mi west of Badger Creek (fig. 3). The Elbert County site is located in

T. 6 S., R. 57 W., sec. 8 and lies immediately west of Beaver Creek (fig. 4).

Sampling Methods for Crops

The sampling protocol for crops was designed to determine the average composition of the crop in each of the six 20-acre fields. For the Elbert County site, whole wheat plants were collected about one month prior to harvest. For this collection, stainless-steel pruning shears were used to cut the plants about 3 inches above the ground. Approximately 30 subsamples of wheat plants were collected randomly in each of the three 20-acre fields. One month later, immediately before harvest, samples of the mature wheat grain were collected. Stainless steel pruning shears were used to cut the plant about 2 inches below the seed head. Again, about 30 subsamples were collected at random in each 20-acre field.

The wheat crop at the Arapahoe County site failed and was plowed under. The only wheat that remained was in a few scattered parts of each 20-acre field; the north control field was almost devoid of wheat plants. Most of the few plants that were still growing in the north control field were removed as whole-plant samples. Few wheat plants remained to produce seed heads. Samples of whole plants and seed heads were collected, if possible, from these limited areas of the fields. About 30 wheat subsamples were collected from the areas where wheat remained in each field. Approximately the western one-half of the Arapahoe County site was replanted with millet. At maturity (September 2000), seed heads were collected from millet in each of the three 20-acre fields. Stainless-steel pruning shears were used to cut the millet plant about 2 inches below the seed head. About 30 millet subsamples were randomly collected throughout each of the three 20-acre fields.

Analytical Methods for Crops

Whole wheat-plant samples (hereinafter referred to as "wheat-plant samples") were dried under forced air at room temperature and were then ground to less than 2 mm using a standard Wiley mill. A split of the dried and ground wheat sample was ashed in a forced-air muffle furnace at 450°C. The ashed aliquot was analyzed for cadmium, copper, molybdenum, nickel, lead, and zinc by using a standard four-acid digestion followed by detection with an inductively-coupledplasma mass spectrometer (Briggs and Meier, 1999). The ash values were then converted to a dry weight basis by using the ash conversion factor. The dried, unashed material was analyzed for arsenic and selenium by using a standard wet oxidation digestion followed by hydride-generation atomic absorption spectrometry (Hageman and Welsch, 1996). Mercury concentration of the dried, unashed material was determined by cold-vapor generation atomic absorption spectrometry after digestion in nitric acid – dichromate solution (O'Leary and others, 1996). Gross alpha activity, gross beta activity,

and plutonium concentration were determined by standard radiological counting methods (Greenberg, 1992; Whittaker and Grothaus, 1979; Lyon, 1980).

The grain samples were dried under forced air at room temperature. For wheat samples, the grain was separated from the head by using a test plot thrashing machine at Colorado State University. The grain was cleaned using forced air and sieving. For millet samples, the grain was separated from the head by placing the entire sample in a heavy plastic bag and manipulating the heads by hand to free the grain. The grain was cleaned using forced air and sieving. The clean samples of wheat and millet grain were then ground to a flour using a commercial table-top mill. A split of each ground sample was ashed in a forced-air muffle furnace at 450°C. The ashed grain samples and the dried, unashed grain samples were subjected to the same analytical protocol discussed previously for ashed and unashed whole-wheat samples.

Quality Assurance for Crops

The accuracy of the crop analysis was ensured by the analysis of NIST SRM 2709, an agricultural soil. Separate splits of this SRM were randomly placed among the crop and soil samples and submitted to the laboratories. If the analytical results for the constituent of interest in the SRM are within an acceptable range, analytical results for the crop samples are accepted. Separate splits of each composited crop sample are analyzed independently and the results averaged to determine the concentration reported for a given constituent.

Crop Data

Wheat-plant samples were collected from the Arapahoe and Elbert County sites on June 7, 2000. Mature wheat-grain samples (seed heads) were collected from each site on July 6, 2000. Millet-grain samples were collected from the Arapahoe County site on September 27, 2000. The trace-element and radioactivity data for the crops are listed in tables 5-7 (in the Data Section at the back of the report).

Discussion of Crop Data

Because most of the Arapahoe County site was plowed under, it was impossible to collect a sample that was representative of the entire field. Therefore, the wheat-plant and wheat-grain data for the Arapahoe County site will have questionable value for monitoring purposes. Again, because of the limited area of the Arapahoe County site that was planted in millet, the millet-grain samples may not be representative of the entire site, and their value for monitoring purposes is questionable. The sample data, however, will help establish baseline and background values for wheat and millet for the Arapahoe County site.

Ground Water

Applications of pesticides, herbicides, or fertilizers (including biosolids) to the land surface can affect the quality of shallow ground water directly by contaminated recharge or by infiltration through contaminated soils or sediments (remobilization). These applications also can affect the quality of shallow ground water indirectly by tilling (which could mobilize subsurface constituents) or by contributions to natural processes such as nitrification. Further, discharge from contaminated alluvial ground water could contaminate surface water (ponds or streams) or bedrock water-supply aquifers. For this report, alluvial ground water is defined as the water contained in subsurface, unconsolidated (uncemented), windor water-transported sediments in current or historical stream channels or flood plains. Bedrock ground water is defined as the water contained in the fractures or pore spaces of the rock (consolidated sediments) that underlies soil or other uncemented materials; the primary bedrock aquifer in the study area is the Laramie-Fox Hills aguifer (Robson and Banta, 1995). Alluvial and bedrock ground water are separate components in the monitoring program but are combined in this report because the data were collected in the same way and the types of data included are the same.

Objectives of Monitoring Ground Water

Ground water was monitored to characterize the hydrology and water quality of the aquifers; to determine if concentrations of nitrate, arsenic, cadmium, copper, chromium, lead, mercury, molybdenum, nickel, selenium, zinc, and plutonium and gross alpha and gross beta activity in the ground water are significantly greater than regulatory standards; and to determine if concentrations of these constituents are increasing with time in ground water at or near the MWRD properties.

Approach for Monitoring Ground Water

Structure maps of the top and base of the Laramie-Fox Hills aquifer were compiled for the study area by using available information such as geophysical logs from oil and gas exploration and other data. The structure maps are included in an interpretive USGS report by Yager and Arnold (2003). The structure maps were used to determine locations for well pairs: two bedrock-aquifer wells and two alluvial-aquifer wells that constitute the recharge-evaluation sites. Multiple wells in the same location enable different zones of ground water to be monitored without having to consider spatial variability and can enable inferences about vertical directions of ground-water flow between zones. Each of the two bedrock-aquifer wells are nested, which means each borehole has two separate piezometers screened at two separate zones. Therefore, three different aquifer zones are monitored at each of the two recharge-evalu-

ation sites in Muddy Creek downgradient from the MWRD properties (fig. 1). Lithologic and well-completion information for these wells is included in the 1999 data report (Stevens and others, 2003). An additional well, DTX11, was drilled in January 2000 to provide additional information about a deeper coarse-grained part of the Muddy Creek alluvial aquifer than is monitored by well DTX9. Completion information for well DTX11 is provided in figure 5 (in the Data Section at the back of the report). In 2000, electronic data-logger (EDL) equipment was installed to continuously monitor precipitation and water levels in wells DTX9, DTX10, and DTX11, and to provide more detailed information about ground-water recharge at that location. These EDL data are included in this report but are not available on the Internet.

Monitoring wells for the expanded monitoring program include selected wells installed as part of the previous monitoring program and new wells. Of the 33 USGS ground-water monitoring wells from the previous study on the MWRD central property, 7 are included in this study (all 7 wells were monitored for water levels, and water-quality samples were collected from 6 of these wells). "D"-numbered wells were drilled before 1999 as part of the previous monitoring program, and "DTX"-numbered wells were drilled in 1999 (fig. 1). Lithologic and well-completion information for these well locations is provided by Stevens and others (2003).

Water levels in the monitoring wells were measured monthly. Data-collection platforms (DCP's) with various sensors were installed during summer and fall of 1999 at three alluvial-aquifer wells (D25, DTX2, and DTX5) to continuously monitor ground-water levels, water temperature, precipitation, and air temperature. The data were transmitted to Denver by satellite and were available on the Internet. The data provided information about the hydrology in the study area and the response of ground water to climate variables.

Water samples were collected from alluvial-aquifer wells on the MWRD properties (fig. 1), and water samples were collected from the shallowest zones of the bedrock aguifer at three locations that are important to alluvial/bedrock groundwater interactions. The remaining USGS monitoring wells were used to provide hydrologic information only. Water samples were collected and analyzed quarterly for physical properties, dissolved major ions and trace elements, and dissolved and total nutrients. Analyses were done by the USGS National Water-Quality Laboratory (NWQL) and included nitrate, arsenic, cadmium, copper, chromium, lead, mercury, molybdenum, nickel, selenium, and zinc. Samples also were collected and analyzed annually for total plutonium. Gross alpha and gross beta activity analyses were included in the 1999 monitoring but were discontinued because of matrix-interference problems caused by the high concentrations of dissolved solids in the ground-water samples. The plutonium and gross alpha and gross beta activity analyses were included in response to public concerns that radioactivity in biosolids could increase from the transfer of water from the Lowry Landfill Superfund site. Water levels and field measurements such as pH and specific conductance were recorded with the collection of each groundwater sample. Blank and replicate samples were analyzed to evaluate bias and variability of the ground-water data. All water-quality data are maintained in a USGS database, and selected data were published in the "USGS Expanded Monitoring Program Near Deer Trail" quarterly reports. Selected water-quality data will be statistically analyzed each year of the program and after about 5 years to determine (1) if concentrations in the ground water are significantly greater than regulatory standards and (2) if the concentrations are increasing significantly with time.

Site Selection for Monitoring Ground Water

Shallow aquifers can be recharged by runoff and streamflow or can contribute water to streamflow and ponds. Therefore, the sites for alluvial-aquifer wells were selected by the USGS according to the following criteria: (1) locations in proximity to a stream channel that could carry runoff from MWRD biosolids-applied fields, (2) locations at the most downstream point of the drainage basin, (3) locations at MWRD property boundaries to represent the condition of ground water leaving the properties and to consider only those effects from activities on MWRD properties and not from other landowners, (4) locations where most of the upstream basin is on MWRD property, (5) locations that represent the larger drainage basins, (6) locations where USGS monitoring wells already existed and where data already had been collected, and (7) locations accessible year round for drilling and sampling wells. Alluvial-aquifer wells were not installed upgradient from MWRD property boundaries because the constituents of concern generally are not conservative along the ground-water flow path; that is, subtracting upgradient concentrations from downgradient concentrations may not represent the effects of biosolids on the ground water for these constituents. Monitoring alluvial ground water near Rattlesnake Creek was a low priority because most of the basin is upstream from the MWRD properties, and that part of the basin that receives biosolids is relatively small. Therefore, the USGS installed two alluvial-aquifer wells on the MWRD north property and four on the MWRD south property (fig. 1); all wells on the MWRD central property used for this study (fig. 1) were installed before 1999 as part of the previous monitoring program.

Bedrock aquifers can be recharged by alluvial ground water or can be a source of water to alluvial aquifers. Therefore, the sites for bedrock-aquifer wells were selected by the USGS according to the following criteria: (1) locations where a particular sandstone sequence within the Laramie-Fox Hills aquifer is present at substantial areal extent and thickness, (2) locations on MWRD property where the bedrock aquifer is present without an alluvial aquifer, (3) locations where the bedrock aquifer is present beneath an alluvial aquifer that could be affected by the application of biosolids, (4) locations where USGS monitoring wells already existed and where data already had been collected, and (5) locations accessible year

round for drilling and sampling wells. Locations where the particular sandstone sequence within the bedrock aquifer is present in this area were determined by the USGS on the basis of the USGS structure map (completed in 1999) of the base of the Laramie-Fox Hills aguifer in this area. For the expanded monitoring program, the USGS installed two new bedrockaquifer wells (DTX8 and DTX10) in 1999, along with corresponding alluvial-aquifer wells (DTX7 and DTX9) (fig. 1), at locations where the bedrock aquifer is present beneath the Muddy Creek alluvial aguifer; the Muddy Creek alluvial aguifer could be affected by the application of biosolids. One previously installed USGS ground-water monitoring well, D29 (fig. 1), was included in this monitoring program because the well is on MWRD property where the bedrock aquifer is present without an alluvial aquifer, and prior sampling data are available.

DCP sites provided information about the variability in space and time of climate and hydrology in the study area as well as about the hydrologic responses to climate. This monitoring program includes three DCP sites, one on each of the MWRD north, south, and central properties (wells DTX2, DTX5, and D25, respectively). The locations of these DCP sites were selected according to the following criteria: (1) locations where alluvial-aquifer wells are sampled, (2) locations near possible streambed-sediment sampling drainages (to indicate likely runoff conditions), (3) locations near other wells so the information may apply to more than one well, (4) locations far enough apart from each other to indicate spatial variability in hydrology, (5) locations needing additional hydrologic information to explain chemical variability (well D25), and (6) locations accessible year round.

Sampling Methods for Ground Water

All data-collection methods used in 2000 were the same as the 1999 methods, which are detailed in the 1999 data report (Stevens and others, 2003). Monthly water-level measurements were made using a vinyl-coated electric tape. DCP and EDL data were automatically recorded hourly. Water-quality samples were collected quarterly using standard USGS methods (Horowitz and others, 1994; Wilde and others, 1998-99). All sampling equipment was used exclusively by the USGS and was used only in the study area to prevent cross contamination from other sites in other study areas. All samples and sampling equipment were kept at all times in the custody of the USGS in locked, guarded facilities.

Analytical Methods for Ground Water

Ground-water samples were submitted to the USGS NWQL in Denver. Most analyses were done by the NWQL. Plutonium analyses were done by Severn Trent Laboratory in Richland, Wash., under a contract with the USGS NWQL; all other analyses were done at the NWQL. The methods used to

analyze the ground-water samples are listed in table 8 along with laboratory minimum reporting levels (MRL) for the elements of interest and minimum detectable concentrations (MDC) for the radiochemical samples.

Quality Assurance for Ground Water

Quality-assurance procedures were implemented during the course of the monitoring program to ensure the quality of the data. Procedures were implemented for water-level measurements, DCP-data and core-data collection, ground-water-sampling preparation, field-parameter measurements, ground-water sampling, and laboratory analysis. Quality-assurance procedures are detailed in the 1999 data report (Stevens and others, 2003).

Ground-Water Data

Monitoring at ground-water sites during 2000 produced meteorologic, hydrologic, and water-quality data. Meteorologic data include precipitation and air temperature at three sites. Hydrogeologic data include the lithologic description of the cores and a well-completion diagram for the new well drilled in 2000, DTX11. Hydrologic data include monthly water levels at all wells and hourly water levels and water temperature at three DCP sites. Water-quality data include analytical results from quarterly sampling.

Meteorologic Data

Precipitation and air temperature were recorded hourly during 2000 at wells D25, DTX2, and DTX5 (figs. 6 through 8 in the Data Section at the back of the report). Precipitation was recorded hourly during April-December 2000 near well DTX11 (fig. 9 in the Data Section at the back of the report). Rain occurred during thunderstorms and frontal storms, usually during April-October. The precipitation data recorded during October through December could represent melted snow. The data indicate air temperatures can fluctuate more than 20°C during the day but were similar for all monitored sites.

Hydrogeologic Data

Well DTX11 was cored by the USGS during drilling in 2000 to provide information about ground water and geology at the new well location. The lithologic description for the well is listed in table 9 (at the back of the report). The details of DTX11 well construction are given in the well-completion diagram shown in figure 5. Lithologic descriptions and well-completion information for the other monitoring wells are provided in the 1999 data report (Stevens and others, 2003).

Well information also is summarized in table 10 (at the back of the report). These details, which include the depth of well, screened interval, materials used, and stickup (fig. 5), provide a physical context for the other ground-water data, such as water levels, and should be considered when comparing data for different wells.

Hydrologic Data

Monthly water-level data and continuous water-level and water-temperature data can be useful for describing the hydrology of the aquifers in the area near Deer Trail. The monthly water-level data for the USGS monitoring wells used in this study are listed in table 11 (at the back of the report), and the continuous water-level and water-temperature data for the three DCP sites (D25, DTX2, and DTX5; fig. 1) are shown in figures 6–8 (at the back of the report). Continuous water-level data for the EDL site (wells DTX9, DTX10, and DTX11; fig. 1) and precipitation data are shown in figure 9 (at the back of the report). Monthly water-level data are listed as depth to water below measuring point at a specific time. Continuous water-level data are shown as daily maximum depth to water below land surface. Water-level data can indicate seasonal effects and can aid in the interpretation of chemical data.

Water-level data also can indicate ground-water recharge information. Robson and others (1981) showed that recharge of the Laramie-Fox Hills aquifer along the margin of the Denver Basin (such as in the Deer Trail area) can be from deeper parts of the Denver Basin, from alluvial aquifers and surficial features, or from infiltration of precipitation on or near outcrop areas. Recharge of the alluvial aquifers in the Deer Trail area can be from the Laramie-Fox Hills aguifer, from surface-water features, or from infiltration of precipitation (Robson and others, 1981). Hydrologic interactions between alluvial and bedrock aquifers can be inferred using water-level data for the same point in time for wells drilled into the aguifers at the same site. The direction of the vertical movement of ground water, or the recharge direction, may be indicated by noting that water moves from areas of high hydraulic head (high water-level elevation) to areas of low hydraulic head (low water-level elevation). For the expanded monitoring program, such interactions were monitored at two recharge-evaluation sites, each of which included one alluvial-aquifer well and one bedrock-aquifer well. The interactions in the bedrock aquifer were monitored in two different zones (designated by "A" [shallow] or "B" [deep] after the well name). Therefore, three aguifer zones (one alluvial, one shallow bedrock, and one deep bedrock) were monitored at the north recharge-evaluation site (wells DTX7 and DTX8), and four aquifer zones (two alluvial, one shallow bedrock, and one deep bedrock) were monitored at the south recharge-evaluation site (wells DTX9, DTX10, and DTX11). Water levels for the paired alluvial-aquifer and bedrock-aquifer wells at the two recharge-evaluation sites are shown in figure 10 (at the back of the report) for the north recharge-evaluation site at discrete

time periods, and in figure 11 (at the back of the report) for the south recharge-evaluation site continuously for the year. The July water levels shown in figure 10 represent hot, dry summer conditions; the August and September water levels shown in figure 10 represent summer-recharge (rainy) conditions.

Water-Quality Data

Water-quality data for samples collected quarterly from 11 alluvial-aquifer and 3 bedrock-aquifer wells (fig. 1) in 2000 are listed in tables 12 and 13 (at the back of the report). Data are given for field parameters, physical properties, major ions, nutrients, and trace elements. In addition, radioactivity data are given for January and October 2000. Quality-control water-quality data for the blank samples are listed in table 14 (at the back of the report), and comparison data for replicate samples are listed in table 15 (at the back of the report).

Discussion of Ground-Water Data

Concentrations for the blank samples (table 14) generally indicate little or no contamination bias. January data indicate a possible slight contamination bias in aluminum, and April data indicate a possible slight contamination bias in nitrogen and zinc from field processing or laboratory analyses. Blank samples were collected April 21, 2000, using the submersible pump, and these data indicate a possible slight contamination bias in concentrations of calcium, magnesium, copper, iron, strontium, and zinc (table 14). This pump is used only at wells DTX8, DTX10, and D29, so only samples from these wells may be affected. Most concentrations for the blank samples were much less than those for the ground-water samples. Data for rerun samples (not included in this report) indicate slight contamination bias in laboratory analyses of aluminum, arsenic, copper, nickel, and zinc. Therefore, the sample data that indicate small concentrations of aluminum, arsenic, copper, nickel, and zinc could be affected by a slight high bias.

The relative percent differences (RPD) between the ground-water samples and the replicate samples were computed to summarize sample variability (table 15). Many of the larger RPD's are due to values or concentrations near the MRL where precision is expected to be poor. In these cases, concentrations may vary little but result in large RPD's. For example, a ground-water sample concentration of 0.01 mg/L and a replicate-sample concentration of 0.02 mg/L would result in an RPD of 67 percent, but the difference might be considered to be within the precision of the method at that concentration. Data values for individual replicate pairs also are listed in table 15 to help determine if large RPD's are the result of substantial differences between replicate-sample concentrations or just small differences between small concentrations. Some RPD's for potassium and nickel, however, likely are large because a few of these concentrations are affected by

analytical bias (table 13). The replicate-sample data indicate generally reproducible analytical results.

The radioactivity data in table 13 are reported in the uncensored form, as received from the laboratory, rather than censored by either the contract or calculated minimum detectable concentration (MDC). Relative to the censored form (data reported as less than the MDC), the uncensored form provides more information about the uncertainty and the very small plutonium concentrations. The negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day. Radioactivity data are produced from instruments that detect radioactive decay (disintegrations) in a sample as counts per minute. The background counts were subtracted from the sample counts, and the resulting value was converted to activity-concentration units of picocuries per liter. Gross alpha and gross beta activity could not be analyzed in the ground-water samples because of interference effects related to the ionic strength of these samples.

In general, the expanded monitoring program is too new for sufficient data to have been collected for meaningful interpretation of the ground-water data. However, the data included in this report indicate alluvial- and bedrock-aquifer hydrology and chemistry are variable in space (from site to site) and in time (from one data-collection time to the next at the same site) in the study area. Water-quality data in this report for samples before July 2000 provide baseline information for concerns about possible contamination of the study area from the Lowry Landfill Superfund site water transfer; no water from the Lowry site was transferred to MWRD until July 2000.

The USGS was asked to evaluate the water-quality data for biosolids effects each year of the monitoring program. Therefore, the results for selected constituents were tested to determine if statistical evidence indicated exceedance of regulatory standards and a monotonic trend in concentration with time.

Regulatory Standards

Regulatory standards that might be used as guidelines to evaluate the ground-water quality in the study area are the human health standards and agricultural standards enforced by the State of Colorado (Colorado Department of Public Health and Environment, 1997). For this report, a one-tailed Sign Test (Helsel and Hirsch, 1995) was used to indicate the level of statistical evidence that selected median constituent concentrations were significantly greater than regulatory standards. The hypotheses tested were as follows:

H_o = median concentration is less than or equal to the regulatory standard, and

 H_a = median concentration is greater than the regulatory standard.

A small p-value result from the Sign Test indicates H_a, the null hypothesis, should be rejected. The confidence level in rejecting H₂ and, therefore, accepting H₂ can be determined by subtracting the p-value from 1 and multiplying by 100. The confidence level also can be thought of as the probability (in percent) that the regulatory standard has been exceeded by the median concentration (table 16 at the back of the report). For example, if the Sign Test for a constituent results in a p-value of 0.100, there is a 90-percent confidence level or probability that the median concentration for that constituent is greater than the regulatory standard. The results of the statistical testing of the data for the 11 alluvial-aquifer wells and 3 bedrockaquifer wells for exceedance of regulatory standards for 11 constituents identified as important by the public are listed in table 16. Nitrate concentrations at well D6 were the only data that had a statistically significant probability that the median concentration exceeded the regulatory standard.

Because quarterly samples have been collected for only 2 years to date, the power of the statistical test (level of statistical evidence) is low. As sampling continues, the power of the test will increase. Radioactivity data were not statistically tested because few radiochemical samples, an insufficient number of samples for statistical testing, were collected at each well since 1999. The distribution of concentrations at each well for selected constituents during 2000 and the regulatory standards used to test the ground-water data are shown in figure 12 (at the back of the report). Time-series graphs (concentration plotted with time) of selected constituents for selected wells are included as figure 13 (at the back of the report).

Trends

Upward monotonic trends in concentration mean that concentration for a parameter increases over time, although not necessarily in a straight line. These upward trends could indicate biosolids, other farm practices, grazing, or even natural processes such as geochemical dissolution are affecting ground water. For this report, the Kendall's tau statistic (Helsel and Hirsch, 1995) was used as an indicator of monotonic correlation between concentration and time. Kendall's tau is a number between negative one and positive one where values approaching negative or positive one indicate increasing strength of the correlation and a number approaching zero indicates decreasing strength of correlation. Positive values of Kendall's tau indicate upward trends, and negative values indicate downward trends. The results of the statistical testing of the data for monotonic trends in 11 constituents identified as important by the public are listed in table 17 (at the back of the report). A p-value is listed to indicate the level of significance of the coefficient, the tau value. The p-value must be less than or equal to 0.05 for tau to be significant with at least 95-percent confidence. Values in table 17 indicating statistically significant trends are shown in bold. Radioactivity data were not statistically tested because only three radiochemical samples, which is an insufficient number of samples for

statistical testing, have been collected at each well since the program began in 1999.

If all data for a constituent were less than the MRL, no trend could be identified (such as for lead at any of the wells). If at least one concentration value for a constituent at a well was greater than the MRL, the data were analyzed for trend. Many of the reported concentrations included in the trend analysis were less than the MRL, and the MRL varied for most of the constituents. Therefore, the trends evaluated by the statistical analyses may be artificial--that is, the difference in MRL, not differences in concentration, may result in an apparent trend. However, none of the statistically significant trends (table 17, values in bold type) were caused by changes in the MRL. The results of trend analysis on the 1999 plus 2000 combined data set (table 17) commonly were different than the results of trend analysis on only the 1999 data set (Stevens and others, 2003). Some of the trend directions changed from upward to downward trends or from downward to upward trends when the 1999 data were retested with the 2000 data, and significance of the trend result increased. Some of these changes may be caused by the low power of the test when few data are used in the test (as in 1999), by changes in the MRL as mentioned previously, or because seasonal effects have not been considered (too few data). Because quarterly samples have been collected for only 2 years to date, the power of the trend test is still low, and too few data are available to consider seasonal effects. As sampling continues, the power of the test will increase and the amount of data available will enable seasonal effects to be removed. Time-series graphs (concentration plotted with time) of selected constituents for selected wells are included as figure 13 (at the back of the report).

Streambed Sediment

Animal waste related to grazing livestock and applications of pesticides, herbicides, or fertilizers (including biosolids) to the land surface could affect surface-water quality directly by contaminated inflow or by runoff over contaminated soil or sediment (remobilization). These applications also can affect surface-water quality indirectly by tilling that mobilizes or changes surface constituents or surface characteristics; by inflow, base flow, or recharge to surface water from contaminated ground water; or by contributions to natural processes such as nitrification. Contaminated surface water could contaminate downstream, previously uncontaminated ponds, streams, streambed sediments, alluvial aquifers, or bedrock water-supply aquifers in aquifer-recharge zones.

Surface-water contamination from biosolids applications is a public concern. However, because streams flow off the MWRD properties only after intense thunderstorms, surface-water sampling is impractical, and monitoring extreme surface-water events is difficult. Monitoring streambed-sediment chemistry is more practical and cost effective and offers greater opportunity to establish comparison or baseline sites

than monitoring surface-water chemistry. For the expanded monitoring program, streambed sediment is defined as the fine-grained alluvium freshly deposited in the drainage bottoms by surface-water flow after rainstorms.

Sediment affected by the application of biosolids could be transported off MWRD property into streambeds when precipitation is intense enough to cause overland flow. Therefore, streambed-sediment chemistry is used as an indirect indicator of surface-water quality because of the close contact between surface flows and sediment during transport. Contaminants in the streambed sediment could cause contamination in ground water or surface water if the contaminants are resuspended in water or leached from the bed sediment. Furthermore, concentrations of trace elements and plutonium and gross alpha and gross beta activity may be higher in the bed sediment than in the surface water.

Objectives of Monitoring Streambed Sediment

Streambed sediments were monitored to determine if concentrations of nitrate, arsenic, cadmium, chromium, copper, lead, mercury, molybdenum, nickel, selenium, zinc, organic carbon, gross alpha and beta radioactivity, and plutonium in sediments derived from (or transported through) biosolids-application areas are significantly higher than in bed sediments derived from nearby farmland that did not receive biosolids applications.

Approach for Monitoring Streambed Sediment

Two small drainage basins were selected for comparison of streambed-sediment chemistry (fig. 1). The basins have similar physical characteristics, but one basin (known as the biosolids-applied basin) receives biosolids applications and is part of the MWRD farming program, and the other basin (known as the control basin) receives no biosolids applications and is farmed privately.

A DCP site (with a rain gage) is near the biosolids basin. The DCP data were transmitted remotely to enable the USGS to determine when rainfall of sufficient intensity and duration had occurred near the basins. For sampling to take place, sufficient rainfall was needed to cause the fine-grained materials and dissolved constituents to wash off the hillsides of the basins into downgradient streambeds.

Paired streambed-sediment samples were collected when freshly deposited streambed sediment was available from both the biosolids basin and the control basin at the same time (after the same storm). The USGS was able to collect only one paired sample during 2000 despite several attempts. The USGS determined that at least 0.5 inch of rain in a single hour had to be received by the tipping-bucket rain gage at DTX2 to cause sufficient runoff in the biosolids-applied basin that a streambed-sediment sample could be collected. This rainfall scenario occurred only once in 2000.

Streambed-sediment samples were analyzed for nitrate, phosphorus, aluminum, arsenic, cadmium, chromium, copper, lead, mercury, molybdenum, nickel, selenium, zinc, carbon, gross alpha and beta radioactivity, and plutonium. The radioactivity analyses were included in response to public concerns that biosolids radioactivity concentrations could increase from the transfer of water from the Lowry Landfill Superfund site. When sufficient storms occur, three to four paired samples per year are planned. When enough paired samples are collected, data will be statistically tested to determine if streambed-sediment chemistry is significantly different between the two basins.

Site Selection for Monitoring Streambed Sediment

Several pairs of similar small basins (one on MWRD property and one on nearby private property) were considered by the USGS for monitoring. Only small (less than 5-mi²) basins were considered because (1) the large biosolids basins did not have a nearby corresponding control basin that had similar characteristics, (2) the thunderstorms likely to cause runoff were usually localized and were not likely to affect two large basins with the same duration and intensity, (3) sampling is more efficient in small basins because each streambed-sediment sample is a composite of sediment collected throughout the basin, and (4) the large basins in the study area are more variable with respect to geology, soil type, and land use. Sediment collected from the large basins is more likely to be affected by this variability and, therefore, may not indicate effects from biosolids. The criteria used to pair basins included distance between basins; proximity to a USGS ground-watermonitoring well that included DCP instrumentation; land use (farmed); likely accessibility even after severe thunderstorms; and similar bedrock geology, soil type, aspect, stream order, channel length, channel slope, relief, and ponding. From the three basin pairs identified as candidates for streambedsediment monitoring (Stevens and others, 2003), the pair on Badger Creek tributaries (on and near the MWRD's north property) was selected. The selected basins are shown as streambed-sediment sampling areas in figure 1.

Sampling Methods for Streambed Sediment

Before sampling, equipment was field washed with phosphate-free detergent, rinsed three times with deionized (DI) water and wrapped in clean plastic bags for transport to the next site. Trace-element sampling equipment received an additional rinse with 5-percent trace-element-grade nitric-acid solution and three more rinses with DI water. Clean sieve cloth was used for each site. The trace-element equipment was allowed to air dry and was stored in plastic bags until use. Stainless-steel equipment (which was not used to prepare trace-element samples) was allowed to air dry and was

wrapped in aluminum foil and stored in sealed plastic containers.

DCP data transmitted by satellite to the USGS were monitored daily throughout the year to determine the occurrence, intensity, and duration of rainfall in the study area. If sufficient rainfall in the area of the paired basins was indicated by the data, the sampling crew visited the sites to determine if the rainfall had produced sediment transport from the hillsides to the alluvial channel in both basins. If transport occurred in both basins, a streambed-sediment sample was collected from the newly transported sediment in the main-stream channel of the basin.

Freshly deposited bed sediment from the basins (fig. 1) was sampled after rainfall runoff by using dedicated equipment for each basin in accordance with the procedures of Radtke (1997). Bed sediment was collected from the alluvial channel in each of the selected basins by using Teflon spoons. The upper 2 cm of fine-grained, wet sediment that collects in depositional areas was removed, placed in a clean sealable plastic bag, and transported in a cooler containing ice to the USGS District laboratory in Denver for processing. Processing consisted of homogenizing, sieving, and bottling the sample. For the trace-element sample, the sediment was washed into an acid-rinsed polypropylene sample jar through a nylon, 0.63 µm sieve using DI water. The sediment in the jar was allowed to settle, and then the clear water was removed from the top of the sample with a plastic syringe. This procedure was repeated until sufficient sediment (about 35 g total) was sieved for the sample. The procedure was repeated using a 2-mm stainless-steel sieve to fill sample containers for analyses of inorganic and organic carbon (500 g in a 1-L, baked glass jar), nutrients (20 g in a polypropylene jar), gross alpha (1 g in a polypropylene jar), and plutonium (150 g in a polypropylene jar). The samples were labeled and transported to the USGS NWQL for analyses. Laboratory analyses consisted of nutrients (total nitrogen and total phosphorus), trace elements (aluminum, arsenic, cadmium, chromium, copper, lead, mercury, molybdenum, nickel, selenium, and zinc), and plutonium and gross alpha and gross beta activity.

Analytical Methods for Streambed Sediment

Trace-element samples were analyzed at the USGS NWQL. Gross alpha, gross beta, and plutonium analyses were done by Severn Trent Laboratory in Richland, Wash., under a contract with the USGS NWQL. The analytical methods used to analyze the streambed-sediment samples and the laboratory MRL's or MDC's are listed in table 18 (at the back of the report).

Quality Assurance for Streambed Sediment

Quality-assurance procedures were implemented during the course of the monitoring program to ensure the quality of the data. The DI water used in cleaning and sample processing was monitored for purity according to procedures given in Horowitz and others (1994). Laboratory and field cleaning procedures were rigorous and designed to prevent contamination of samples. Before sample collection, all sampling equipment and materials were cleaned according to standard procedures given in Horowitz and others (1994), Radtke (1997), and Wilde and others (1998-99).

If results for a particular constituent were questionable, the sample was reanalyzed by the laboratory. If results from the second analysis were more consistent with known characteristics of the site or the particular sample, the new results were used instead of the previous results; otherwise, the initial result was retained. The analytical quality-assurance practices and procedures of the NWQL are described in Friedman and Erdmann (1982).

Sufficient streambed sediment was available for a replicate sample, so a replicate July 17 sample of the control site also was analyzed. These data (table 19 at the back of the report) provide information about variability in the sediment itself or contributed by field conditions, sampling equipment, or the laboratory. Other quality-control samples for 2000 included a laboratory replicate, a laboratory method blank, and a laboratory spike analyzed only for plutonium and gross alpha and gross beta activity. The laboratory replicate was used to assess analytical precision. Laboratory blanks were used to assess contamination bias. The laboratory spike was prepared by the laboratory by using known concentrations of a constituent to assess recovery and analytical precision. The quality-control samples provided information about the bias and variability contributed by the laboratory but not the bias and variability contributed by field conditions or sampling equipment or about the variability of the sediment.

Streambed-Sediment Data

The USGS collected a single paired sample July 17, 2000. Data for these two samples and the replicate are included in table 19 at the back of the report.

Discussion of Streambed-Sediment Data

A comparison of the limited sample data with the replicate data indicates that variability within the sample is about the same as variability between the basins for this single paired sample. However, a statistical evaluation is not useful for the limited data collected.

The radioactivity data are reported in the uncensored form as received from the laboratory rather than censored by either the contract or calculated MDC. Relative to the censored form (data reported as less than the MDC), the uncensored form provides more information about the uncertainty,

the very small concentrations of plutonium, and the gross alpha and gross beta activity. The negative activity concentration reported for one of the radiochemical samples means the sample count was less than the laboratory background count for that day. Radioactivity data are produced from instruments that detect radioactive decay (disintegrations) in a sample as counts per minute. The background count was subtracted from the sample count, and the resulting value was converted to activity-concentration units of picocuries per gram.

The streambed-sediment data in this report should provide baseline information for concerns about contamination of the study area from the Lowry Landfill Superfund site water transfer. No water from the Lowry site was transferred to MWRD until July 25, 2000, after the July 17 streambed-sediment sample was collected (table 1 and fig. 2).

REFERENCES CITED

- Briggs, P.H., 1996, Forty elements by inductively coupled plasma-atomic emission spectrometry, *in* Arbogast, B.F., ed., 1996, Analytical methods manual for the Mineral Resource Surveys Program, U.S. Geological Survey: U.S. Geological Survey Open-File Report 96–525, p. 77–94.
- Briggs, P.H., and Meier, A.L., 1999, The determination of forty-two elements in geological materials by inductively coupled plasma-mass spectrometry: U.S. Geological Survey Open-File Report 99–166, 15 p.
- Colorado Department of Public Health and Environment, 1997, Basic standards for ground water: 5CCR 1002–41, July 14, 1997, 56 p.
- 1998, Biosolids regulation: 5CCR 1002–64, January 12, 1998, 53 p.
- Drever, J.I., 1988, The geochemistry of natural waters (2d ed.): New Jersey, Prentice-Hall, p. 328–329.
- Drew, L.J., Schuenemeyer, J.H., and Bawiec, W.J., 1979, Petroleum exhaustion maps of the Cretaceous "D–J" sandstone stratigraphic interval of the Denver Basin: U.S. Geological Survey Miscellaneous Investigations Series Map I–1138, scale 1:200,000, sheet 2.
- Faires, L.M., 1993, Methods of analysis by the U.S. Geological Survey National Water Quality Laboratory--Determination of metals in water by inductively coupled plasma-mass spectrometry: U.S. Geological Survey Open-File Report 92–634, 28 p.
- Fishman, M.J., ed., 1993, Methods of analysis by the U.S. Geological Survey National Water Quality Laboratory-Determination of inorganic and Organic constituents in water and fluvial sediments: U.S. Geological Survey Open-File Report 93–125, 217 p.

- Fishman, M.J., and Friedman, L.C., 1989, Methods for determination of inorganic substances in water and fluvial sediments: U.S. Geological Survey Techniques of Water-Resources Investigations, book 5, chap. A1, 545 p.
- Friedman, L.C., and Erdmann, D.E., 1982, Quality-assurance practices for the chemical and biological analyses of water and fluvial sediments: U.S. Geological Survey Techniques of Water-Resources Investigations, book 5, chap. A6, 181 p.
- Greenberg, A.E., ed., 1992, Method 7110—Gross alpha and gross beta radioactivity (total, suspended, and dissolved), *in* American Public Health Association and the AWWA, Standard Methods for the Examination of Water and Wastewater (18th ed.): Washington, D.C., p. 7–1 through 7–14.
- Hageman, P.L., and Welsch, Eric, 1996, Arsenic, antimony, and selenium by flow injection or continuous flow-hydride generation-atomic absorption spectrometry, *in* Arbogast, B.F., ed., Analytical methods manual for the Mineral Resource Surveys Program, U.S. Geological Survey: U.S. Geological Survey Open-File Report 96–525, p. 24–30.
- Helsel, D.R., and Hirsch, R.M., 1995, Statistical methods in water resources (3d ed.): Amsterdam, Elsevier Studies in Environmental Science 49, 529 p.
- Horowitz, A.J., Demas, C.R., Fitzgerald, K.K., Miller, T.L., and Rickert, D.A., 1994, U.S. Geological Survey protocol for the collection and processing of surface-water samples for the subsequent determination of inorganic constituents in filtered water: U.S. Geological Survey Open-File Report 94–539, 57 p.
- Jones, S.R., and Garbarino, J.R., 1999, Methods of analysis by the U.S. Geological Survey National Water Quality Laboratory-Determination of arsenic and selenium in water and sediment by graphite furnace-atomic absorption spectrometry: U.S. Geological Survey Open-File Report 98–639, 39 p.
- Larsen, L.S., Baber, T.G., Wesswick, E.L., McCoy, D.E., andHarman, J.B., 1966, Soil survey of Elbert County, Colorado:U.S. Department of Agriculture Soil Conservation ServiceSoil Survey series, 79 p.
- Larsen, L.S., and Brown, J.B., 1971, Soil survey of ArapahoeCounty, Colorado: U.S. Department of Agriculture SoilConservation Service Soil Survey series, 78 p.
- Lyon, W.S., ed., 1980, Radioelement analysis—Progress and problems: Ann Arbor Science, p. 215–221 and 223–230.
- Major, T.J., Robson, S.G., Romero, J.C., and Zawistowski, Stanley, 1983, Hydrogeologic data from parts of the Denver Basin, Colorado: U.S. Geological Survey Open-File Report 83–274, 425 p.

- McLain, Betty, 1993, Methods of analysis by the U.S. Geological Survey National Water Quality Laboratory--Determination of chromium in water by graphite furnace atomic absorption spectrophotometry: U.S. Geological Survey Open-File Report 93–449, 16 p.
- Motooka, Jerry, 1996, Organometallic halide extraction for 10 elements by inductively coupled plasma-atomic emission spectrometry, *in* Arbogast, B.F., ed., Analytical methods manual for the Mineral Resource Surveys Program, U.S. Geological Survey: U.S. Geological Survey Open-File Report 96–525, p. 102–108.
- O'Leary, R.M., Hageman, P.L., and Crock, J.G., 1996, Mercury in water, geologic, and plant materials by continuous flow-cold vapor-atomic absorption spectrometry, *in* Arbogast, B.F., ed., Analytical methods manual for the Mineral Resource Surveys Program, U.S. Geological Survey: U.S. Geological Survey Open-File Report 96–525, p. 42–50.
- Patton, C.J., and Truitt, E.P., 1992, Methods of analysis by the U.S. Geological Survey National Water Quality Laboratory-Determination of total phosphorus by a Kjeldahl digestion method and an automated colorimetric finish that includes dialysis: U.S. Geological Survey Open-File Report 92–146, 39 p.
- Radtke, D.B., 1997, National field manual for the collection of water-quality data—Chapter A8. Bottom-material samples:
 U.S. Geological Survey Techniques of Water-Resources Investigations, Book 9, Chapter A8, 52 p.
- Robson, S.G., and Banta, E.R, 1995, Ground water atlas of the United States, segment 2, Arizona, Colorado, New Mexico, Utah: U.S. Geological Survey Hydrologic Investigations Atlas 730–C, p. C20–C22.
- Robson, S.G., Wacinski, Andrew, Zawistowski, Stanley, and Romero, J.C., 1981, Geologic structure, hydrology, and water quality of the Laramie-Fox Hills aquifer in the Denver Basin, Colorado: U.S. Geological Survey Hydrologic Investigations Atlas Map HA–650, scale 1:500,000, sheet 1.
- Seaber, P.R., Kapinos, F.P., and Knapp, G.L., 1987, Hydrologic unit maps: U.S. Geological Survey Water-Supply Paper 2294, 63 p.
- Sharps, J.A., 1980, Geologic map of the Limon 1 x 2 quadrangle, Colorado and Kansas: U.S. Geological Survey Miscellaneous Investigations Series Map I–1250, scale 1:250,000, sheet 1.
- Stevens, M.R., Yager, T.J.B., Smith, D.B., and Crock, J.G., 2003, Biosolids, soils, ground-water, and streambed-sediment data for a biosolids-application area near Deer Trail, Colorado: U.S. Geological Survey Open-File Report 02–51, 118 p.

- Struzeski, T.M., DeGiacomo, W.J., and Zayhowski, E.J., 1996, Methods of analysis by the U.S. Geological Survey National Water Quality Laboratory--Determination of dissolved aluminum and boron in water by inductively coupled plasmatomic emission spectrometry: U.S. Geological Survey Open-File Report 96–149, 17 p.
- U.S. Geological Survey, 1974, Hydrologic unit map—1974, State of Colorado: U.S. Geological Survey Hydrologic Unit Map 1974, scale 1:500,000, 1 sheet.
- 1980, Land use and land cover, 1975, Limon, Colorado; Kansas: U.S. Geological Survey Land Use Series L–191, scale 1:250,000, 1 sheet.
- Whittaker, E.L., and Grothaus, G.E., 1979, Acid dissolution method for the analysis of plutonium in soil, EPA–600/7–79–081, Environmental Monitoring and Support Laboratory, Office of Research and Development: Las Vegas, U.S. Environmental Protection Agency, p. 23–59.
- Wilde, F.D., Radtke, D.B., Gibs, J., and Iwatsubo, R.T., 1998–1999, National field manual for the Collection of water-quality data—Chapters A1-A5: U.S. Geological Survey Techniques of Water-Resources Investigations, Book 9, Chapters A1, A2, A3, A4, and A5, variously paginated.
- Yager, T.J.B., and Arnold, L.R., 2003, Hydrogeology of a biosolids-application site near Deer Trail, Colorado, 1993–99:
 U.S. Geological Survey Water Resources Investigations
 Report 03–4209, 90 p.

Data Section

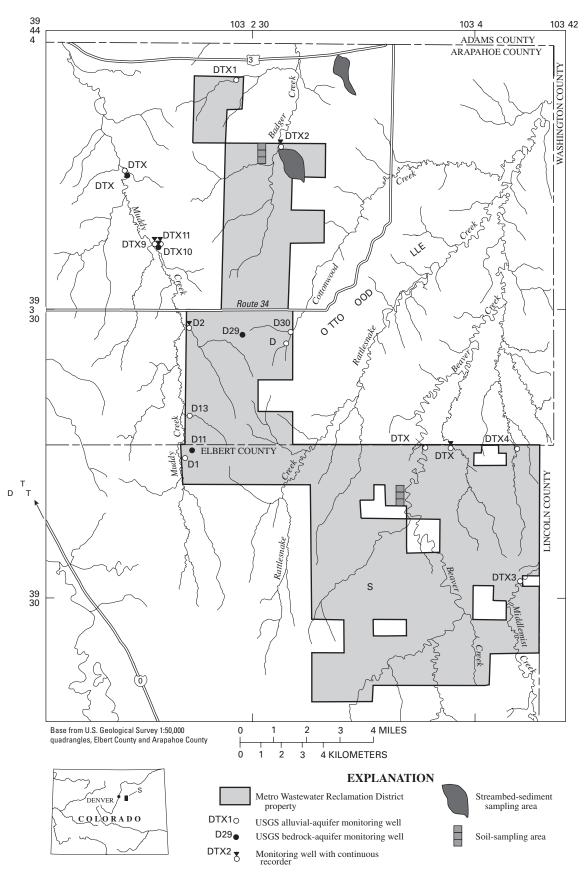


Figure 1. Location of study area and USGS monitorings sites near Deer Trail, Colorado, 2000. MWRD property boundaries shown are from 1999.

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Figure 2 Metro Wastewater Reclamation District biosolids-application areas (METROGRO Farm) near Deer Trail, Colorado, 2000 (from Metro Wastewater Reclamation District).

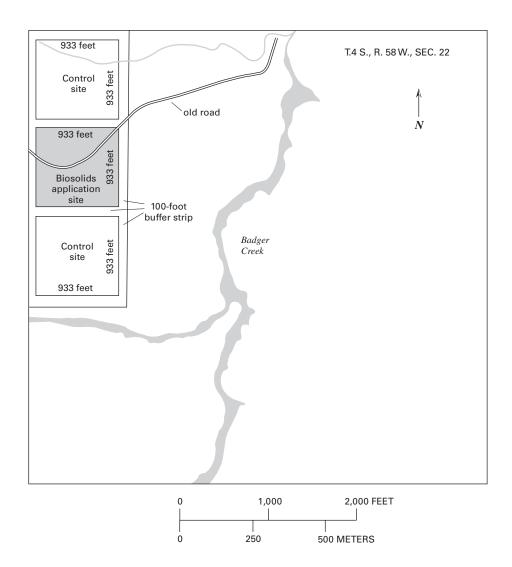


Figure 3. Arapahoe County, Colorado, soil-monitoring site: T. 4 S., R. 58 W., sec. 22 (from Metro Wastewater Reclamation District).



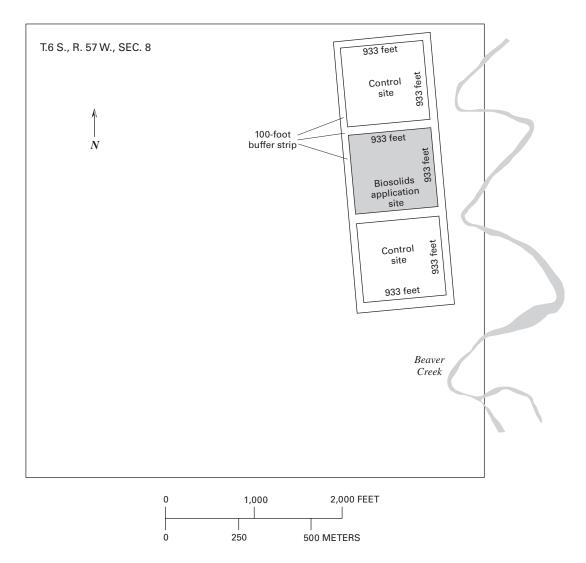


Figure 4. Elbert County, Colorado, soil-monitoring site: T. 6 S., R. 57 W., sec. 8 (from Metro Wastewater Reclamation District)..

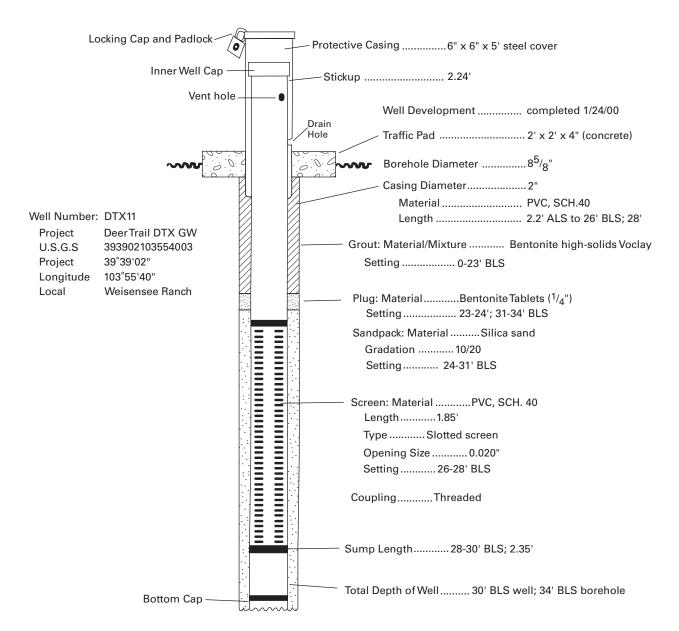
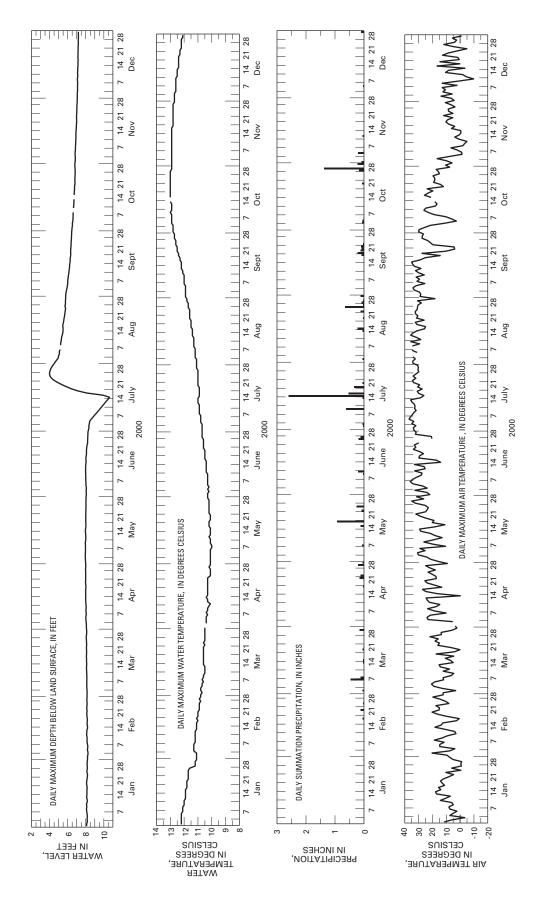


Figure 5. Well-completion information for U.S. Geological Survey monitoring well DTX11 near Deer Trail, Colorado, 2000 (PVC, polyvinyl chloride; SCH., schedule; ALS, above land surface; BLS, below land surface).



Continuous water-level, water-temperature, precipitation, and air-temperature data for well D25 near Deer Trail, Colorado, 2000

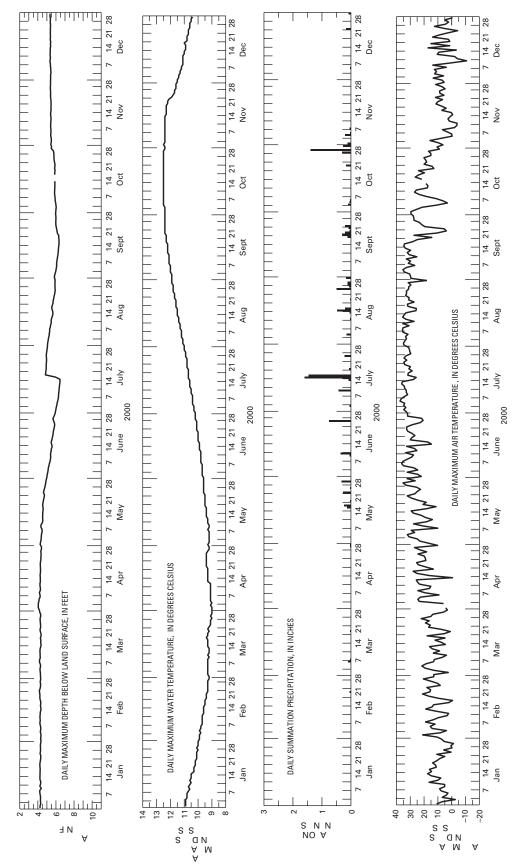
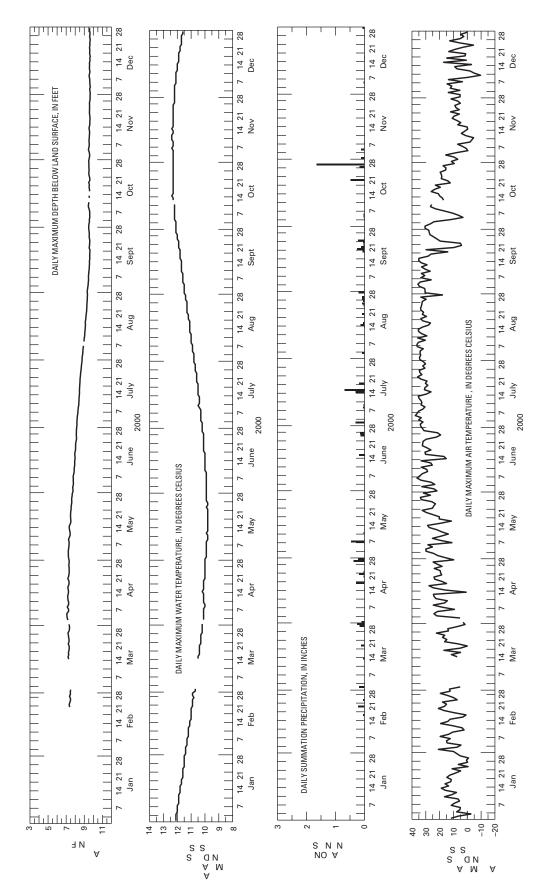
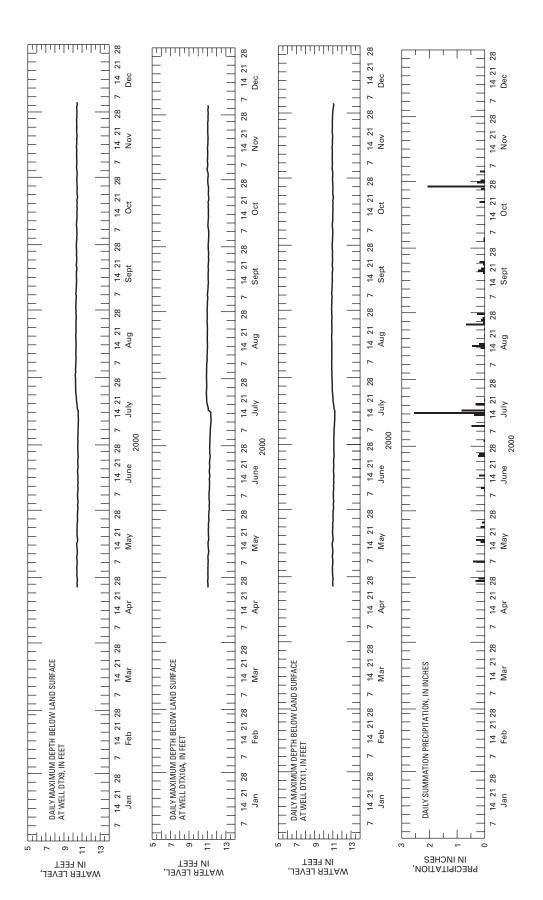


Figure 7. Continuous water-level, water-temperature, precipitation, and air-temperature data for well DTX2 near Deer Trail, Colorado, 2000



Continuous water-level, water-temperature, precipitation, and air-temperature data for well DTX5 near Deer Trail, Colorado, 2000



Continuous water-level data for wells DTX9, DTX10A, and DTX11, and precipitation data for the EDL recharge-evaluation site near Deer Trail, Colorado, Figure 9. 2000.

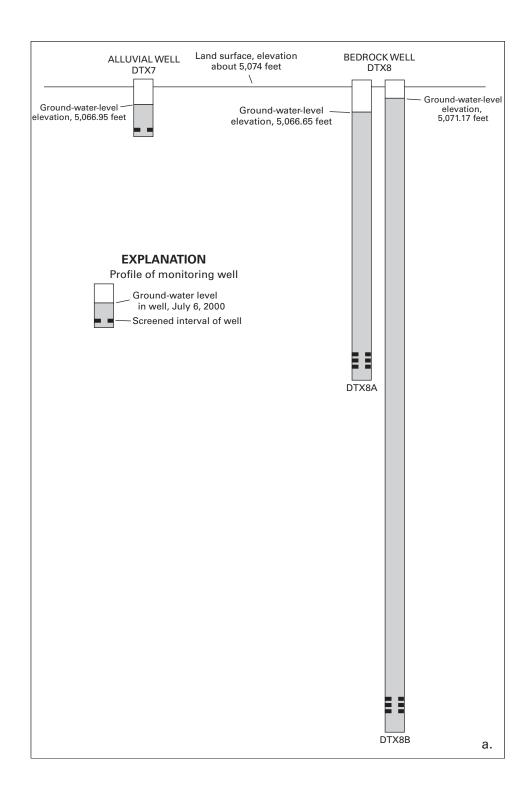


Figure 10. Water levels for the recharge-evaluation site containing well DTX7 and nested well DTX8 (includes DTX8A and DTX8B) near Deer Trail, Colorado, for (a) July, (b) August, and (c) September 2000. (Elevation is in feet above NAVD 88.)

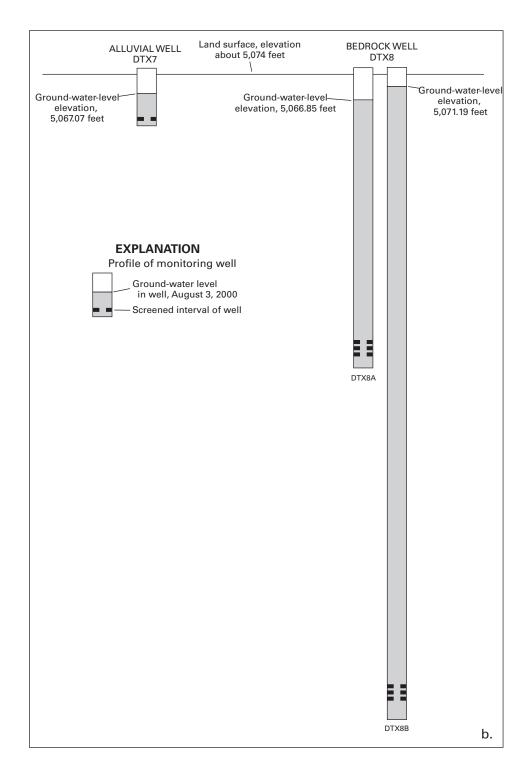


Figure 10. Water levels for the recharge-evaluation site containing well DTX7 and nested well DTX8 (includes DTX8A and DTX8B) near Deer Trail, Colorado, for (a) July, (b) August, and (c) September 2000. (Elevation is in feet above NAVD 88.)—Continued.

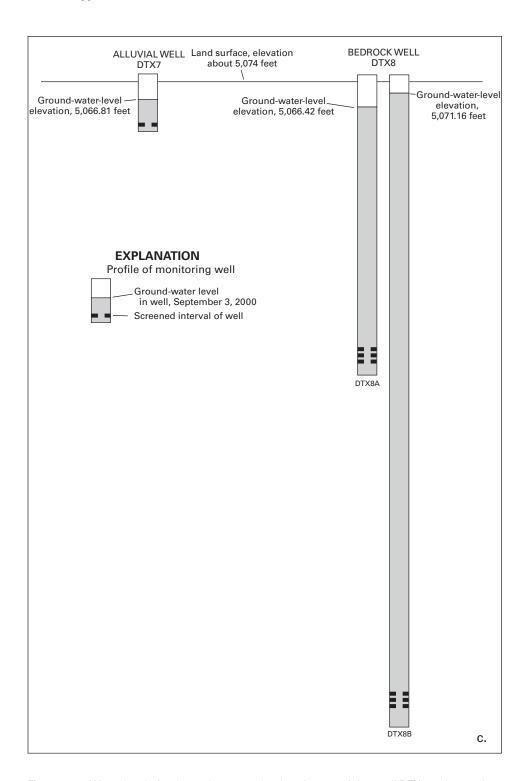


Figure 10. Water levels for the recharge-evaluation site containing well DTX7 and nested well DTX8 (includes DTX8A and DTX8B) near Deer Trail, Colorado, for (a) July, (b) August, and (c) September 2000. (Elevation is in feet above NAVD 88.)—Continued.

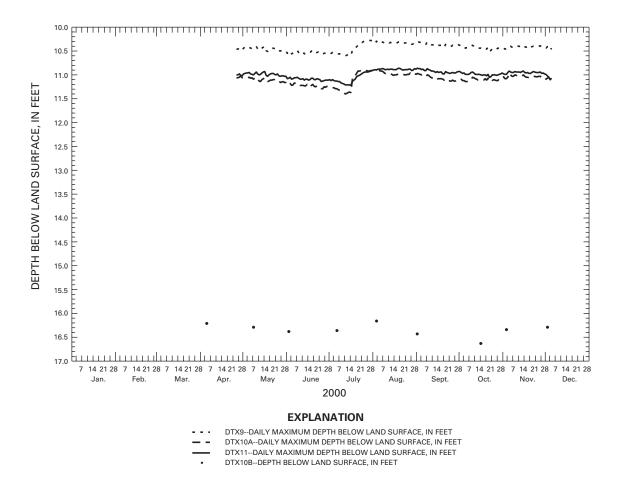


Figure 11. Water levels for the recharge-evaluation site containing wells DTX9, DTX10, and DTX11 near Deer Trail, Colorado, 2000.

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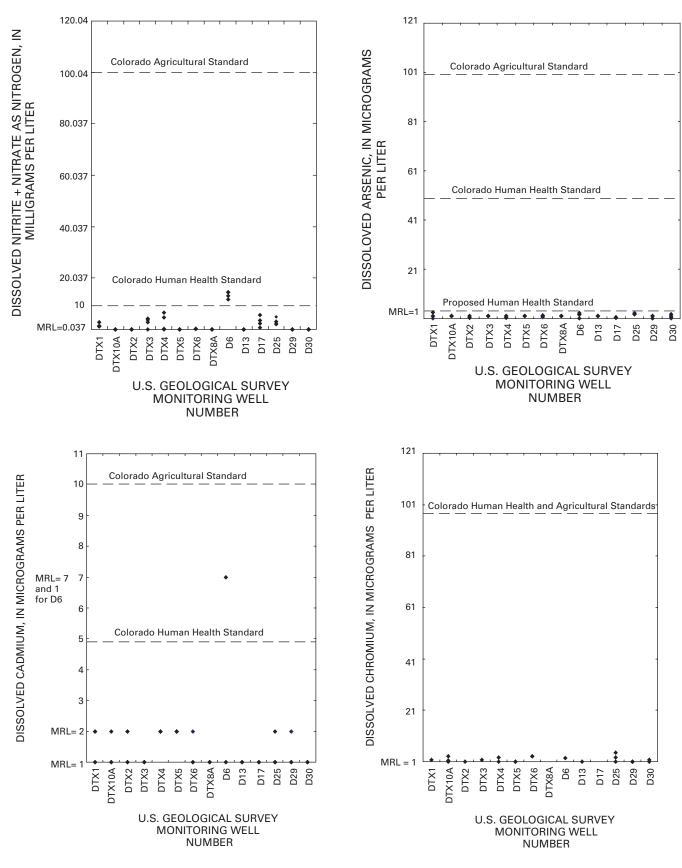


Figure 12. Distribution of ground-water constituent data collected near Deer Trail, Colorado, compared to regulatory standards for selected constituents, 2000.

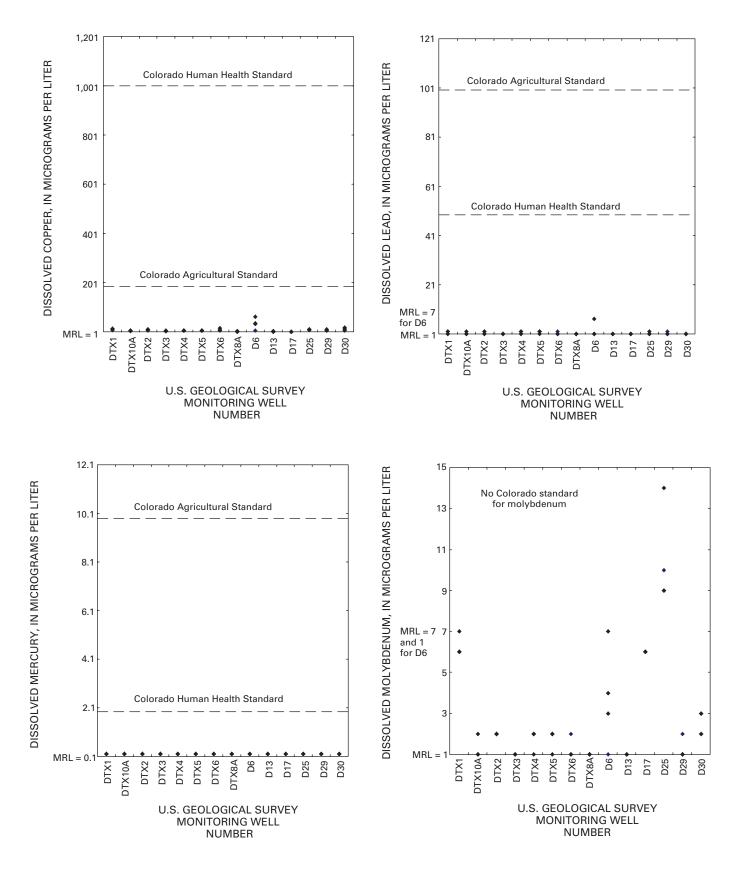


Figure 12. Distribution of ground-water constituent data collected near Deer Trail, Colorado, compared to regulatory standards for selected constituents, 2000—Continued.

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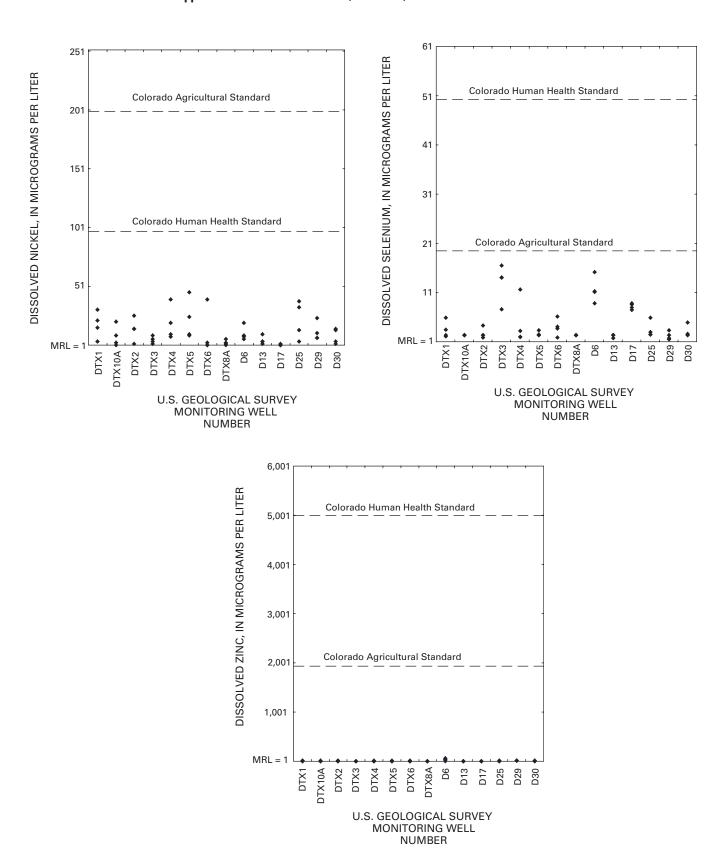
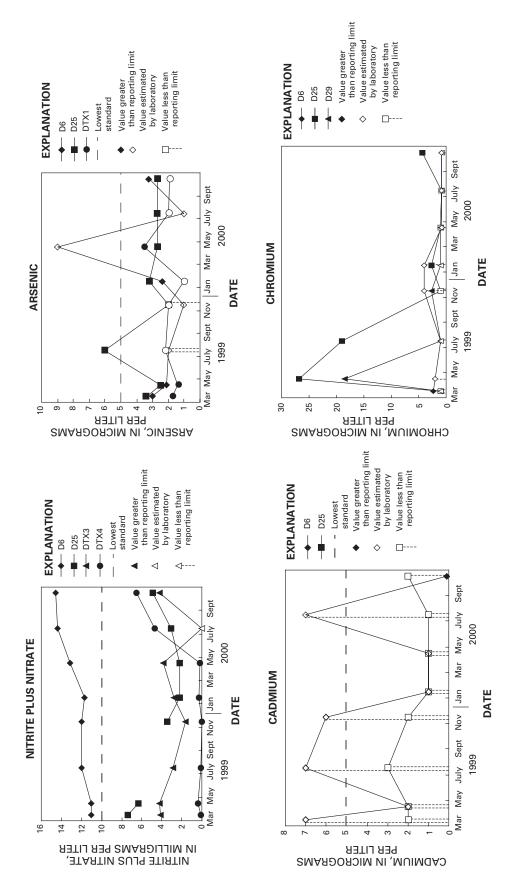


Figure 12. Distribution of ground-water constituent data collected near Deer Trail, Colorado, compared to regulatory standards for selected constituents, 2000—Continued.



Ground-water concentrations near Deer Trail, Colorado, for selected constituents and selected wells, 1999-2000. Figure 13.

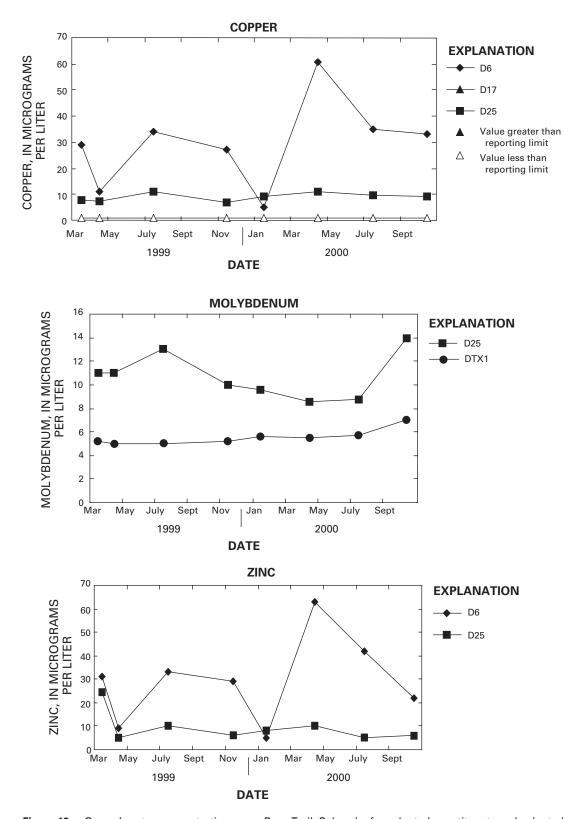


Figure 13. Ground-water concentrations near Deer Trail, Colorado, for selected constituents and selected wells, 1999-2000—Continued.

TERMS AND ABBREVIATIONS

The following terms and abbreviations are used in tables 1-19:

bls below land surface

bmp below the measuring point of the well casing

mi mile ft feet

hhmm 24-hour time

mm/dd/yy numerical date format for two-digit month/two-digit day/ and the last two

digits of the year

C Celsius
cm centimeter
degree
in. or " inch

μS/cm microsiemens per centimeter at 25°Celsius

pH units are the negative base-10 log of the hydrogen-ion activity in moles per liter

mg/L milligrams per liter µg/L micrograms per liter pCi/L picocuries per liter

dissolved refers to that fraction of material in a water sample that passes through a

0.45-µm membrane filter

> greater than < less than

g/kg grams per kilogram mg/kg milligrams per kilogram pCi/g picocuries per gram µg/g micrograms per gram

NAD 83 North American Datum 1983

NAVD 88 North American Vertical Datum 1988

 Table 1.
 Biosolids applications near Deer Trail, Colorado, 2000

[All information provided by Metro Wastewater Reclamation District; DC, destination code (shown in fig. 2); legal description is of the form quarter-section township range; Ha, hectares; CAKE, Grade I Class B biosolids; MAC, biosolids amended with wood fiber; dT, dry tons; dMT, dry metric tons; lbs/Acre, pounds per acre; Kg/Ha, kilograms per hectare; Y, yes; N, No; O/GR, oats and grass; --, no data]

2		Area applied to	pplied 5	ć	Biosolids	Start	Stop	Total	Total	Total c	Total dry tons	Loading rate, tons per acre	g rate, r acre	Nitrogen loading rate	gen g rate	d	Recla-
2	Legal description	Acres	На	- County	product applied	date	date	loads	tons	Τρ	dMT	Cake dT	dMT	lbs/Acre	Кд/На	d S	mation project
300	N 1/2 SEC 15 T5S R58W	160	64.8	ARAPAHOE	CAKE	7/18/00	7/21/00	82	1,835	299	271.25	1.87	4.19	46	51.57	WHEAT	z
300	N 1/2 SEC 15 T5S R58W	160	64.8	ARAPAHOE	CAKE	9/18/00	11/18/00	51	1,121	180	163.30	1.13	2.53	28	31.39	WHEAT	Z
301	S 1/2 SEC 15 T5S R58W	155.7	63.0	ARAPAHOE	CAKE	7/22/00	7/25/00	95	2,125	329	298.47	2.11	4.73	52	58.30	WHEAT	z
301	S 1/2 SEC 15 T5S R58W	78	31.6	ARAPAHOE	CAKE/MAC	9/19/00	11/18/00	45	666	165	149.69	1.92	4.30	50	56.06	WHEAT	z
303	S 1/2 SEC 16 T5S R58W	148	59.9	ARAPAHOE	CAKE/MAC	7/25/00	7/30/00	118	2,628	424	384.65	2.77	6.21	70	78.48	WHEAT	z
306	N 1/2 SEC 20 T5S R58W	, 107.3	43.4	ARAPAHOE	CAKE	3/10/00	3/17/00	45	1,028	164	148.78	1.53	3.43	43	48.21	WHEAT	z
307	S 1/2 SEC 20 T5S R58W	107.1	43.3	ARAPAHOE	CAKE	3/11/00	6/21/00	80	1,824	279	253.11	2.61	5.85	69	77.36	WHEAT	z
308	N 1/2 SEC 21 T5S R58W	156.6	63.4	ARAPAHOE	CAKE/MAC	11/22/00	11/29/00	93	2,090	350	317.52	2.04	4.57	28	65.02	WHEAT	z
309	S 1/2 SEC 21 T5S R58W	129.3	52.3	ARAPAHOE	CAKE	11/23/00	11/28/00	75	1,707	268	243.13	2.07	4.64	99	62.78	WHEAT	z
310	N 1/2 SEC 22 T5S R58W	69.3	28.0	ARAPAHOE	CAKE	11/19/00	11/22/00	54	1,219	274	248.57	2.74	6.14	29	75.11	WHEAT	z
311	S 1/2 SEC 22 T5S R58W	66.1	26.8	ARAPAHOE	CAKE/MAC	11/18/00	11/21/00	48	1,081	180	163.30	2.50	5.61	99	73.99	WHEAT	z
312	N 1/2 SEC 28 T5S R58W	, 166.3	67.3	ARAPAHOE	CAKE/MAC	11/29/00	12/5/00	135	2,970	510	462.67	2.72	6.10	83	93.05	WHEAT	z
313	S 1/2 SEC 28 T5S R58W	224.7	6.06	ARAPAHOE	CAKE	11/30/00	12/8/00	71	1,585	251	227.71	1.12	2.51	32	35.88	WHEAT	z
315	S 1/2 SEC 29 T5S R58W	23.9	6.7	ARAPAHOE	CAKE	12/8/00	12/9/00	18	416	29	80.78	2.80	6.28	92	85.20	WHEAT	z
317	S 1/2 SEC 32 T5S R58W	159.6	64.6	ARAPAHOE	CAKE/MAC	66/6/9	3/4/00	722	16,122	2,674	2,425.85	16.30	36.55	415	465.26	GRASS	Y
318	N 1/2 SEC 33 T5S R58W	145.3	58.8	ARAPAHOE	CAKE/MAC	12/11/00	12/17/00	116	2,490	278	252.20	2.68	6.01	80	69.68	WHEAT	z
319	S 1/2 SEC 33 T5S R58W	94.3	38.2	ARAPAHOE	CAKE/MAC	66/6/L	2/28/00	137	3,094	519	470.84	5.20	11.66	130	145.74	O/GR	Y

 Table 1.
 Biosolids applications near Deer Trail, Colorado, 2000—Continued

[All information provided by Metro Wastewater Reclamation District; DC, destination code (shown in fig. 2); legal description is of the form quarter-section section township range; Ha, hectares; CAKE, Grade I Class B biosolids; MAC, biosolids amended with wood fiber; dT, dry tons; dMT, dry metric tons; lbs/Acre, pounds per acre; Kg/Ha, kilograms per hectare; Y, yes; N, No; O/GR, oats and grass; --, no data]

2		Area applied to	applied to	ć	Biosolids	Start	Stop	Total	Total	Total d	Total dry tons	Loading rate, tons per acre	g rate, r acre	Nitrogen Ioading rate	gen g rate	į	Recla-
3	regal description	Acres	Е	County	applied	date	date	loads	tons	₽	dMT	Cake dT	dMT	lbs/Acre	Кд/На	9	mation project
319	S 1/2 SEC 33 T5S R58W	50.2	20.3	ARAPAHOE	CAKE/MAC	12/9/00	12/11/00	41	902	157	142.43	2.83	6.35	81	90.81	WHEAT	z
322	N 1/2 SEC 3 T6S R58W	215.7	87.3	ELBERT	CAKE	12/19/00	12/26/00	121	2,760	447	405.52	2.07	4.64	99	72.87	WHEAT	z
323	S 1/2 SEC 3 T6S R58W	230.6	93.3	ELBERT	CAKE	12/26/00	01/04/01	184	4,143	683	619.62	2.96	6.64	;	ı	1	1
324	N 1/2 SEC 4 T6S R58W	38	15.4	ELBERT	CAKE/MAC	12/17/00	12/18/00	29	649	1117	106.14	2.68	6.01	84	94.17	WHEAT	z
325	S 1/2 SEC 4 T6S R58W	38	15.4	ELBERT	CAKE/MAC	12/16/00	12/20/00	31	708	124	112.49	2.87	6.43	68	82.66	WHEAT	z
340	W 1/2 SEC 9 T4S R58W	258.4	104.6	ARAPAHOE	CAKE	10/06/00	06/20/01	66	2,207	487	441.81	1.88	4.22	1	ı	1	1
341	N 1/2 SEC 8 T4S R58W	300	121.4	ARAPAHOE	CAKE/MAC	8/29/00	11/17/00	231	5,018	810	734.83	2.60	5.83	69	77.36	WHEAT	z
342	S 1/2 SEC 8 T4S R58W	311.3	126.0	ARAPAHOE	CAKE/MAC	8/22/00	11/13/00	217	4,763	828	751.16	2.49	5.58	99	73.99	WHEAT	z
343	N 1/2 SEC 17 T4S R58W	327.4	132.5	ARAPAHOE	CAKE/MAC	8/18/00	11/10/00	259	5,777	984	892.68	2.91	6.52	81	90.81	WHEAT	Z
344	S 1/2 SEC 17 T4S R58W	280	113.3	ARAPAHOE	CAKE/MAC	8/8/00	8/18/00	223	5,125	855	775.66	2.91	6.52	75	84.08	WHEAT	Z
345	N 1/2 SEC 21 T4S R58W	310	125.5	ARAPAHOE	CAKE/MAC	7/29/00	11/5/00	255	5,823	975	884.52	2.94	6.59	78	87.45	WHEAT	z
346	S 1/2 SEC 21 T4S R58W	300	121.4	ARAPAHOE	CAKE/MAC	7/31/00	10/26/00	224	5,000	831	753.88	2.67	5.99	69	77.36	WHEAT	z
351	N 1/2 SEC 28 T4S R58W	290	117.4	ARAPAHOE	CAKE/MAC	8/27/00	9/16/00	210	4,591	743	674.05	2.38	5.34	59	66.14	WHEAT	z
352	S 1/2 SEC 28 T4S R58W	299	121.0	ARAPAHOE	CAKE	9/2/00	00/6/6	211	4,674	749	679.49	2.51	5.63	61	68.39	WHEAT	z
412	N 1/2 SEC 1 T6S R57W	343	138.8	ELBERT	CAKE	9/26/00	10/2/00	140	3,117	480	435.46	1.40	3.14	40	44.84	WHEAT	z
413	S 1/2 SEC 1 T6S R57W	343	138.8	ELBERT	CAKE/MAC	10/2/00	10/15/00	248	5,465	883	801.06	2.43	5.45	99	73.99	WHEAT	z
425	S 1/2 SEC 11 T6S R57W	20	8.1	ELBERT	CAKE	1/15/00	1/15/00	12	268	39	35.38	1.95	4.37	53	59.42	WHEAT	z
448	N 1/2 SEC 21 T6S R57W	223	90.2	ELBERT	CAKE/MAC	2/10/00	3/26/00	178	3,984	655	594.22	2.81	6.30	77	86.32	WHEAT	z
449	S 1/2 SEC 21 T6S R57W	123.7	50.1	ELBERT	CAKE/MAC	3/4/00	3/30/00	100	2,237	365	331.13	2.74	6.14	92	85.20	WHEAT	z

 Table 1.
 Biosolids applications near Deer Trail, Colorado, 2000—Continued

[All information provided by Metro Wastewater Reclamation District; DC, destination code (shown in fig. 2); legal description is of the form quarter-section section township range; Ha, hectares; CAKE, Grade I Class B biosolids; MAC, biosolids amended with wood fiber; dT, dry tons; dMT, dry metric tons; lbs/Acre, pounds per acre; Kg/Ha, kilograms per hectare; Y, yes; N, No; O/GR, oats and grass; --, no data]

2		Area applied to	pplied o	4	Biosolids	Start	Stop	Total	Total	Total d	Total dry tons	Loading rate, tons per acre	g rate, r acre	Nitrogen Ioading rate	gen g rate	į	Recla-
3	regal description	Acres	Ha	Codility	applied	date	date	loads	tons	Ą	dMT	Cake dT	TMP	lbs/Acre	Кg/На	5	project
450	N 1/2 SEC 22 T6S R57W	219	9.88	ELBERT	CAKE/MAC	2/21/00	3/25/00	169	3,745	618	59.095	2.57	5.76	73	81.84	WHEAT	z
451	S 1/2 SEC 22 T6S R57W	219	88.6	ELBERT	CAKE	2/21/00	3/26/00	93	2,064	327	296.65	1.49	3.34	41	45.97	WHEAT	Z
452	N 1/2 SEC 23 T6S R57W	303	122.6	ELBERT	CAKE/MAC	12/26/99	1/13/00	180	4,014	648	587.87	1.95	4.37	99	62.78	WHEAT	Z
453	S 1/2 SEC 23 T6S R57W	303	122.6	ELBERT	CAKE/MAC	1/7/00	1/19/00	133	2,980	502	455.41	1.43	3.21	42	47.09	WHEAT	Z
454	N 1/2 SEC 24 T6S R57W	30	12.1	ELBERT	CAKE	1/10/00	1/11/00	19	415	49	58.06	2.13	4.78	57	63.90	WHEAT	Z
455	S 1/2 SEC 24 T6S R57W	35	14.2	ELBERT	CAKE	1/19/00	1/21/00	28	630	94	85.28	2.69	6.03	74	82.96	WHEAT	Z
458	N 1/2 SEC 30 T6S R57W	106.2	43.0	ELBERT	CAKE/MAC	1/5/99	9/4/01	317	7,058	1,391	1,261.92	13.10	29.37	1	1	1	1
461	S 1/2 SEC 29 T6S R57W	23.9	6.7	ELBERT	CAKE/MAC	1/9/99	8/24/01	92	1,696	294	266.72	12.30	27.58	1	1	1	1
462	N 1/2 SEC 28 T6S R57W	107	43.3	ELBERT	CAKE/MAC	1/23/00	7/6/00	68	2,003	346	313.89	2.73	6.12	87	97.54	WHEAT	z
463	S 1/2 SEC 28 T6S R57W	54	21.9	ELBERT	CAKE/MAC	1/26/00	1/29/00	43	923	152	137.89	2.56	5.74	73	81.84	WHEAT	z
464	N 1/2 SEC 27 T6S R57W	213	86.2	ELBERT	CAKE/MAC	1/29/00	2/29/00	169	3,737	605	548.86	2.77	6.21	78	87.45	WHEAT	z
465	S 1/2 SEC 27 T6S R57W	172	9.69	ELBERT	CAKE/MAC	1/21/00	1/26/00	138	3,023	466	422.76	2.56	5.74	72	80.72	WHEAT	z
474	N 1/2 SEC 33 T6S R57W	52.2	21.1	ELBERT	CAKE/MAC	2/7/00	2/9/00	31	<i>L</i> 69	119	107.96	1.99	4.46	58	65.02	WHEAT	z
476	N 1/2 SEC 34 T6S R57W	61.6	24.9	ELBERT	CAKE/MAC	2/4/00	2/8/00	50	1,129	181	164.20	2.69	6.03	83	93.05	WHEAT	Z

Table 2. Methods used to analyze biosolids and soil samples collected near Deer Trail, Colorado, 2000

Element or property	Medium	Analytical method	Reference
Arsenic	Soils and biosolids	HG-AAS ¹	Hageman and Welsch (1996)
Cadmium	Biosolids	ICP-MS ²	Briggs and Meier (1999)
Cadmium	Soils	ICP-AES ³	Motooka (1996)
Copper	Biosolids	ICP-MS ²	Briggs and Meier (1999)
Copper	Soils	ICP-AES ³	Briggs (1996)
Lead	Biosolids	ICP-MS ²	Briggs and Meier (1999)
Lead	Soils	ICP-AES ³	Briggs (1996)
Mercury	Soils and biosolids	CV-AAS ⁴	O'Leary and others (1996)
Molybdenum	Biosolids	ICP-MS ²	Briggs and Meier (1999)
Molybdenum	Soils	ICP-AES ³	Motooka (1996)
Nickel	Biosolids	ICP-MS ²	Briggs and Meier (1999)
Nickel	Soils	ICP-AES ³	Briggs (1996)
Selenium	Soils and biosolids	HG-AAS ¹	Hageman and Welsch (1996)
Zinc	Biosolids	ICP-MS ²	Briggs and Meier (1999)
Zinc	Soils	ICP-AES ³	Briggs (1996)
Gross Alpha, Total	Soils and biosolids	Radiological method	Greenberg (1992)
Gross Beta, Total	Soils and biosolids	Radiological method	Greenberg (1992)
Plutonium-238, Total	Soils and biosolids	Radiological method	Whittaker and Grothaus (1979); Lyon (1980)
Plutonium-239+240, Total	Soils and biosolids	Radiological method	Whittaker and Grothaus (1979); Lyon (1980)

¹Hydride Generation-Atomic Absorption Spectrometry.

²Inductively Coupled Plasma-Mass Spectrometry.

³Inductively Coupled Plasma-Atomic Emission Spectrometry.

⁴Continuous Flow-Cold Vapor-Atomic Absorption Spectrometry.

Table 3. Chemical and radioactivity data for biosolids samples collected at the Metro Wastewater Reclamation District during 2000

[ppm, parts per million; pCi/g, picocuries per gram; ±, plus or minus the analytical uncertainty; maximum allowable concentrations from Colorado Department of Public Health and Environment, 1998]

Element or property	Maximum allowable for Grade I	March	June	August	September	October	November	December
Arsenic, ppm	41	1.7	1.8	2.0	2.0	1.8	1.8	1.7
Cadmium, ppm	39	3.0	2.8	2.7	2.9	2.9	5.4	3.6
Copper, ppm	1,500	530	999	580	260	570	570	610
Lead, ppm	300	49	99	78	75	120	70	65
Mercury, ppm	17	1.6	1.6	2.7	1.5	1.6	1.7	1.6
Molybdenum, ppm	No standard set for Grade I, 75 for Grade II	22	26	31	33	31	23	20
Nickel, ppm	420	30	31	33	30	29	27	28
Selenium, ppm	100	8.6	111	12	11	12	11	10
Zinc, ppm	2,800	610	630	700	069	029	069	700
Gross Alpha, pCi/g	40	23 ± 14	44 ± 11	36 ± 12	45 ± 12	43 ± 11	44 ± 11	36 ± 11
Gross Beta, pCi/g	No standard set for Grade I	26±5	23 ± 6	27 ± 5	29 ± 5	29 ± 5	25 ± 4	22 ± 4
Plutonium 238, pCi/g	No standard set for Grade I	0.00 ± 0.01	0.00 ± 0.01	0.02 ± 0.03	0.01 ± 0.02	0.00 ± 0.01	-0.01 ± 0.01	0.01 ± 0.02
Plutonium 239+240, pCi/g No standard set for Grade I	/g No standard set for Grade I	0.00 ± 0.01	0.00 ± 0.01	0.00 ± 0.01	0.00 ± 0.01	-0.01 ± 0.01	0.00 ± 0.01	0.00 ± 0.01

Table 4. Comparison of radioactivity data from two laboratories for biosolids samples [Samples were run at two different laboratories for quality-assurance purposes; concentrations in picocuries per gram]

Constituent or property	Data from Acculabs received in 2001 (reported in table 3)	Data from Severn Trent Laboratory received in 2002 ¹
	June 2000 biosolids sample	
Gross alpha	44 ± 11	40.2 ± 10
Gross beta	23 ± 6	30.1 ± 5.2
Plutonium-238	0.00 ± 0.01	0.00 ± 0.02
Plutonium-239+240	0.00 ± 0.01	0.00 ± 0.00
	NIST Standard Reference Material	2781
Gross alpha	$34 \pm 13, 40 \pm 11, 51 \pm 13, 37 \pm 11, 60 \pm 13, 44 \pm 12, 42 \pm 12$	34 ± 9 , 39 ± 10 , 45 ± 11
Gross beta	$28 \pm 7, 27 \pm 6, 35 \pm 6, 31 \pm 5, 28 \pm 5, 30 \pm 5, 29 \pm 5$	$21 \pm 4, 23 \pm 4, 23 \pm 4$
Plutonium-238	$0.00 \pm 0.01, 0.00 \pm 0.00, 0.00 \pm 0.01, 0.00 \pm 0.01,$ $0.01 \pm 0.02, 0.00 \pm 0.01, 0.01 \pm 0.02$	-0.001 ± 0.002 , 0.00 ± 0.025 , 0.00 ± 0.024
Plutonium-239+240	$0.01 \pm 0.02, 0.00 \pm 0.01, 0.00 \pm 0.01, 0.00 \pm 0.01,$ $0.00 \pm 0.01, 0.00 \pm 0.01, 0.00 \pm 0.00$	$0.001 \pm 0.002, 0.00 \pm 0.027, 0.01 \pm 0.02$

¹ U.S. Geological Survey contract laboratory used for all U.S. Geological Survey water resources radiochemical analyses in 1999 through 2002.

Table 5. Trace-element data for wheat whole-plant samples collected from the soil-monitoring sites near Deer Trail, Colorado, September 2000

[All data in milligrams per kilogram; <, less than]

	Ar	apahoe County sit	te		Elbert County site	
Element	North (control) field	Middle (biosolids application) field	South (control) field	North (control) field	Middle (biosolids application) field	South (control) field
Arsenic	0.12	0.15	0.28	< 0.05	< 0.05	< 0.05
Cadmium	.05	.13	.14	.03	.03	.02
Copper	5.4	5.8	5.4	5.5	4.0	4.0
Lead	.17	.30	.37	.06	.04	.07
Mercury	.006	.005	.008	.006	.007	.006
Molybdenum	.86	.86	.79	1.71	1.06	1.08
Nickel	.94	1.20	1.25	.51	.45	.44
Selenium	.62	.44	.36	1.70	.40	.16
Zinc	17	15	15	21	15	16

Table 6. Trace-element and radioactivity data for wheat-grain samples collected from the soil-monitoring sites near Deer Trail, Colorado, September 2000

 $[mg/kg, milligram\ per\ kilogram; pCi/g, picocuries\ per\ gram; <, less\ than; \pm, plus\ or\ minus\ the\ analytical\ uncertainty]$

	Arapahoe	County site	i	Elbert County site	
Element or property	South (control) field	Middle (biosolids application) field	North (control) field	Middle (biosolids application) field	South (control) field
Arsenic, mg/kg	< 0.05	< 0.05	< 0.05	< 0.05	< 0.05
Cadmium, mg/kg	.03	.04	.02	.02	.02
Copper, mg/kg	5.1	6.6	4.9	4.8	4.5
Lead, mg/kg	.01	.02	<.01	.01	<.01
Mercury, mg/kg	<.004	<.004	<.004	<.004	<.004
Molybdenum, mg/kg	.43	.80	.70	.51	.61
Nickel, mg/kg	.74	1.35	.43	.50	.50
Selenium, mg/kg	.41	1.4	2.1	.64	.38
Zinc, mg/kg	20	20	26	25	22
Gross Alpha, pCi/g			1 ±-1	1 ± -1	1 ± -1
Gross Beta, pCi/g			6 ±-2	$6 \pm - 2$	5 ± -2
Plutonium-238, pCi/g			0.00 ± -0.03	0.00 ± -0.01	0.00 ± -0.01
Plutonium-239+240, pCi/g			0.00 ± -0.00	0.00 ± -0.00	0.00 ± -0.01

Table 7. Trace-element data for millet-grain samples collected from the Arapahoe County soil-monitoring site near Deer Trail, Colorado, September 2000

[in milligrams per kilogram; <, less than]

		Arapahoe County site	
Element	North (control) field	Middle (biosolids application) field	South (control) field
Arsenic	0.07	0.04	0.06
Cadmium	.05	.05	.06
Copper	7.0	7.1	7.4
Lead	.21	.02	.06
Mercury	<.004	<.004	<.004
Molybdenum	.73	.78	.65
Nickel	6.4	6.5	5.3
Selenium	.26	.24	.27
Zinc	22	19	21

Table 8. Methods used to analyze ground-water samples collected near Deer Trail, Colorado, 2000

[MRL, minimum reporting level, and dilutions for samples having high specific conductance may result in higher MRL's for some samples; MDC, minimum detectable concentration (radiochemical samples); ICP, inductively coupled plasma; AA, atomic absorption spectrometry; MS, mass spectroscopy; ASF, automated segmented-flow spectrophotometry; IC, ion chromatography; µS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; µg/L, micrograms per liter; GF, graphite furnace; pCi/L, picocuries per liter]

Property or constituent	Units	Analytical method	MRL or MDC	Method Reference
	Maj	or ions and mineral characteristics		
Specific conductance, lab	μS/cm	Wheatstone bridge	2.6	Fishman and Friedman (1989)
oH, lab	units	Electrometric electrode	0.1	Fishman and Friedman (1989)
Calcium, dissolved	mg/L	ICP	0.02	Fishman (1993)
Magnesium, dissolved	mg/L	ICP	0.014	Fishman (1993)
Sodium, dissolved	mg/L	ICP	0.09	Fishman (1993)
Potassium, dissolved	mg/L	AA	0.24	Fishman and Friedman (1989)
Acid-neutralizing capacity, lab as CaCO ₃	mg/L	Electrometric titration	1	Fishman and Friedman (1989)
Sulfate, dissolved	mg/L	IC	0.31	Fishman and Friedman (1989)
Chloride, dissolved	mg/L	IC	0.29	Fishman and Friedman (1989)
Fluoride, dissolved	mg/L	Colorimetry, ASF, ion-selective electrode	0.1	Fishman and Friedman (1989)
Bromide, dissolved	mg/L	Colorimetry, ASF, fluorescein	0.01	Fishman and Friedman (1989)
Silica, dissolved	mg/L	Colorimetry, ASF, molybdate blue	0.09	Fishman (1993)
Dissolved solids, residue at 180°C	mg/L	Gravimetric	10	Fishman and Friedman (1989)
		Nutrients		
Nitrite plus nitrate, dissolved as N	mg/L	Colorimetry, ASF, cadmium reduc-	0.037	Fishman (1993)
		tion, diazotization		
Nitrogen, ammonia, dissolved as N	mg/L	Colorimetry, ASF, salicylate- hypochlorite	0.029	Fishman (1993)
Nitrogen, ammonia plus organic, total as N	mg/L	Colorimetry, ASF, microkjeldahl digestion	0.1	Unpublished
Nitrogen, ammonia plus organic, dissolved as N	mg/L	Colorimetry, ASF, microkjeldahl digestion	0.1	Patton and Truitt (1992)
Phosphorus, total as P	mg/L	Colorimetry, ASF, microkjeldahl digestion	0.05	Patton and Truitt (1992)
Phosphorus, dissolved as P	mg/L	Colorimetry, ASF, microkjeldahl digestion	0.05	Patton and Truitt (1992)
		Trace elements		
Aluminum, dissolved as Al	ug/L	ICP-MS	1	Faires (1993)
Antimony, dissolved as Sb		ICP-MS	1	Faires (1993)
Arsenic, dissolved as As		Hydride generation	2	Jones and Garbarino (1999)
Barium, dissolved as Ba		ICP-MS	1	Faires (1993)
,	F-0, -		-	

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Table 8. Methods used to analyze ground-water samples collected near Deer Trail, Colorado, 2000—Continued

[MRL, minimum reporting level, and dilutions for samples having high specific conductance may result in higher MRL's for some samples; MDC, minimum detectable concentration (radiochemical samples); ICP, inductively coupled plasma; AA, atomic absorption spectrometry; MS, mass spectroscopy; ASF, automated segmented-flow spectrophotometry; IC, ion chromatography; µS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; µg/L, micrograms per liter; GF, graphite furnace; pCi/L, picocuries per liter]

Property or constituent	Units	Analytical method	MRL or MDC	Method Reference
Boron, dissolved as B	μg/L IC	P-MS	16	Struzeski, DeGiacomo, and
				Zayhowski (1996)
Cadmium, dissolved as Cd	μg/L IC	P-MS	1	Faires (1993)
Chromium, dissolved as Cr	μg/L G	F AA	0.8	McLain (1993)
Cobalt, dissolved as Co	μg/L IC	P-MS	1	Faires (1993)
Copper, dissolved as Cu	μg/L IC	P-MS	1	Faires (1993)
Iron, dissolved as Fe	μg/L IC	P .	3	Fishman (1993)
Lead, dissolved as Pb	μg/L IC	P-MS	1	Faires (1993)
Manganese, dissolved as Mn	μg/L IC	P-MS	1	Faires (1993)
Mercury, dissolved as Hg	μg/L H	ydride generation	0.2	Fishman and Friedman (1989)
Molybdenum, dissolved as Mo	μg/L IC	P-MS	1	Faires (1993)
Nickel, dissolved as Ni	μg/L IC	P-MS	1	Faires (1993)
Selenium, dissolved as Se	μg/L H	ydride generation	2.4	Jones and Garbarino (1999)
Silver, dissolved as Ag	μg/L IC	P-MS	1	Faires (1993)
Strontium, dissolved as Sr	μg/L IC	P-MS	1	Fishman (1993)
Zinc, dissolved as Zn	μg/L IC	P-MS	1	Faires (1993)
		Radioactivity		
Uranium, natural dissolved	μg/L IC	P-MS	1	Faires (1993)
Plutonium-238, total	pCi/L A	lpha spectrometry	0.1*	Not Available
Plutonium-239+240, total	pCi/L A	lpha spectrometry	0.1*	Not Available

^{*}Contract required MDC; reported value may be lower depending upon the sample.

Table 9. Lithologic description for well DTX11 in 2000 near Deer Trail, Colorado

[Description from geologist's notes; cm, centimeter; bottom 6 feet of borehole sloughed with wet, sandy silt before well could be completed]

Depth below land surface	Source of informa- tion	Description of material
0-20 feet	Core	Medium-brown and dark-brown, dry, silty clay grading
		into damp, medium-brown, sandy silt.
20-23 feet	Core	Wet, brown silt and very fine-grained sand.
23-24.3 feet	Core	Dry, hard, orange clay grading into very dark gray clay;
		some small shale pebbles and sandstone fragments.
24.3-28 feet	Core	Very wet, granitic, interbedded coarse sand with pebbles
		and wet, gray, fine-grained sand and silt.
28-38 feet	Core	Wet, mostly gray, fine-grained sand and silt with a few
		thin (less than 1 inch) beds of coarse granitic sand and
		pebbles.
38-40 feet	Core	Dry, crumbly, gray and black striped bedrock (very fine-
		grained sandy silt interbedded with shale) grading into
		dry, crumbly, black shale.

Table 10. Information for U.S. Geological Survey monitoring wells near Deer Trail, Colorado, 2000

[Latitude and longitude are in the format degrees minutes seconds referenced to NAD 83; bmp, below measuring point (stickup); Wells shown in bold are sampled for water quality; Elev., elevation in feet above NAVD 88; Metro, Metro Wastewater Reclamation District]

							Screened Interval	nterval						14/211
Well	Latitude	Latitude Longitude	County	Date drilled	Stickup (feet)	Total depth (feet bmp)	Top (feet bmp)	Bottom (feet bmp)	Screen length (feet)	Screen opening (inches)	Sump length (feet)	Elev. of measuring point (feet)	Property owner	vvell diam- eter (inches)
DTX1	394333	1035251	ARAPAHOE 02/16/99	02/16/99	2.56	25.50	20.59	22.19	1.60	0.010	3.31	4,909	Metro	2
DTX2*	394149	1035138	ARAPAHOE 02/16/99	02/16/99	3.23	20.50	15.59	17.19	1.60	0.010	3.31	4,903	Metro	2
DTX3	393024	1034328	ELBERT	02/12/99	3.11	18.71	13.80	15.40	1.60	0.010	3.31	5,195	Metro	2
DTX4	393358	1034342	ELBERT	02/10/99	2.70	16.72	11.81	13.41	1.60	0.010	3.31	4,957	Metro	2
DTX5*	393358	1034548	ELBERT	02/10/99	2.30	20.90	16.09	17.69	1.60	0.010	3.21	4,975	Metro	2
DTX6	393358	1034648	ELBERT	02/9/99	2.43	39.21	34.30	35.90	1.60	0.010	3.31	4,970	Metro	2
DTX7	394054	1035646	ARAPAHOE 02/18/99	02/18/99	2.77	16.10	11.19	12.79	1.60	0.010	3.31	5,075	Price	2
DTX8A	394054	1035645	ARAPAHOE 03/2/99	03/2/99	2.46	77.52	67.56	71.83	4.27	0.010	5.69	5,076	Price	2
DTX8B	394054	1035645	ARAPAHOE 03/2/99	03/2/99	2.49	177.48	167.52	171.79	4.27	0.010	5.69	5,076	Price	2
DTX9*	393902	1035540	ARAPAHOE 02/17/99	02/17/99	2.46	30.15	22.72	24.32	1.60	0.010	5.83	5,121	Weisensee	2
DTX10A*	393902	1035540	ARAPAHOE 03/4/99	03/4/99	2.03	61.97	52.01	56.28	4.27	0.010	5.69	5,122	Weisensee	2
DTX10B	393902	1035540	ARAPAHOE 03/4/99	03/4/99	2.11	121.73	111.77	116.04	4.27	0.010	5.69	5,122	Weisensee	2
DTX11*	393902	1035540	ARAPAHOE 01/19/00	01/19/00	2.24	32	28	30	1.85	0.020	2.35	5,122	Weisensee	2
9Q	393633	1035122	ARAPAHOE 09/12/93	09/12/93	2.65	25	15	25	10	0.010	0.3	5,128.78	Metro	2
D11a	393345	1035423	ELBERT	10/23/97	2.46	143.03	112.65	122.65	10	0.010	20.38	5,377	Metro	2
D13	393442	1035438	ARAPAHOE 04/4/94	04/4/94	1.81	16	9	16	10	0.010	0.3	5,235.33	Metro	2
D17	393334	1035436	ELBERT	04/5/94	06.0	21	11	21	10	0.010	0.3	5,277.73	Metro	2
D25*	393702	1035442	ARAPAHOE 05/1/95	05/1/95	2.23	23	13	23	10	0.010	0.3	5,167.13	Metro	2
D29	393641	1035248	ARAPAHOE 11/4/97	11/4/97	2.38	183.19	147.81	157.81	10	0.010	25.38	5,371	Metro	2
D30	393655	1035122	ARAPAHOE 05/5/95	05/5/95	1.98	19	6	19	10	0.010	0.3	5,096.43	Metro	2

*Wells having continuous-recorder equipment.

Table 11. Monthly water-level data for U.S. Geological Survey monitoring wells near Deer Trail, Colorado, 2000

[Et1, Solinst electric tape; Et2, Slope Indicator electric tape; St, Steel tape; -, no data; *, continuous-recorder site; W.L. bmp, water level below measuring point, in feet]

Well no.	January (01/12/00) W.L. bmp, Et1	February (02/04/00) W.L. bmp, Et1	March (03/03/00) W.L. bmp, Et1	April (04/05/00) W.L. bmp, Et1	May (05/08/00) W.L. bmp, Et1	June (06/02/00) W.L. bmp, Et1	July (07/06/00) W.L. bmp, Et1	August (08/03/00) W.L. bmp, Et1	August (08/03/00) W.L. bmp, Et2	August (08/03/00) W.L. bmp, St	September (09/01/00) W.L. bmp, Et1	October (10/10/00- 10/18/00) W.L. bmp, Et1	November (11/03/00) W.L. bmp, Et1	December (12/02/00) W.L. bmp, Et1
DTX1	7.87	7.84	7.84	7.53	7.65	7.99	8.90	9.03	ł	ŀ	9.23	08.6	69'6	09.6
DTX2*	7.25	7.31	7.24	7.07	7.39	8.00	10.37	8.13	ł	I	8.76	8.86	8.47	8.40
DTX3	8.90	9.01	9.11	9.05	9.29	9.42	9.62	98.6	9.81	9.83	10.04	10.24	9.82	10.31
DTX4	7.94	8.09	8.05	7.77	7.98	69.6	68.6	11.01	11.02	11.01	11.98	12.51	12.62	12.69
DTX5*	9.74	06.6	88.6	9.45	9.73	10.15	10.26	11.19	ł	ŀ	11.72	11.97	12.00	12.04
DTX6	21.67	21.75	21.74	21.45	21.39	21.56	21.79	21.95	ŀ	ł	22.20	22.49	22.57	22.61
DTX7	7.56	7.52	7.44	7.25	7.31	7.66	8.57	8.45	ł	I	8.71	89.8	8.47	8.30
DTX8A	7.92	7.88	7.75	7.48	7.77	8.42	9.35	9.15	ł	ŀ	9.58	9.58	9.33	8.73
DTX8B	4.78	4.88	4.82	4.63	4.70	4.83	4.88	4.86	ł	I	4.89	4.97	4.82	4.81
DTX9	12.81	12.87	12.85	12.81	12.88	12.96	13.00	12.77	12.74	12.74	12.76	12.89	12.87	12.84
DTX10A	13.00	13.09	13.07	12.97	13.11	13.23	13.31	12.93	ł	I	13.00	13.12	13.09	13.07
DTX10B	18.67	18.71	18.60	18.32	18.40	18.49	18.47	18.27	ŀ	ı	18.54	18.74	18.45	18.40
DTX11*	1	1	13.41	13.37	13.20	13.29	13.35	13.11	ŀ	ı	13.10	13.23	13.20	13.17
D6	9.54	9.61	9.61	9.51	9.18	9.20	9.42	9.31	9.29	ı	9.41	9.54	9.48	9.24
D11a	113.24	113.25	113.25	113.00	113.28	113.30	113.16	113.13	ł	I	113.10	112.82	113.21	113.11
D13	7.46	7.75	7.16	6.63	6.92	7.37	8.08	6.82	6.81	08.9	8.61	8.47	8.19	7.68
D17	10.99	11.00	10.99	10.91	10.74	10.77	11.08	11.52	11.51	11.44	12.41	12.65	12.60	12.48
D25*	10.35	10.37	10.37	10.24	10.19	10.29	10.68	7.13	ŀ	ı	7.98	8.81	90.6	9.27
D29	154.25	154.74	154.52	154.24	154.32	154.67	154.35	154.38	ł	I	154.29	154.19	154.49	154.57
D30	5.19	5.20	5.10	4.64	4.87	5.26	5.98	4.95	4.95	4.94	5.28	5.38	4.52	4.64

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Table 12. Field-parameter data associated with ground-water sampling near Deer Trail, Colorado, 2000 [μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter, --, no data because equipment malfunctioned]

Well	Sample date (mm/dd/yy)	Sample time (hhmm)	Specific conduc- tance, field (µS/cm)	pH, field (stan- dard units)	Water temperature (degrees Celsius)	Water level before sampling, depth below measuring point (feet)	Oxygen, dissolved (mg/L)
DTX1	01/10/00	1345	4,200	7.0	11.1	7.84	0.50
DTX1	04/14/00	1220	4,210	7.0	11.3	7.46	0.52
DTX1	07/07/00	940	4,310	6.7	11.1	8.92	0.62
DTX1	10/17/00	1200	4,060	7.1	12.3	9.80	0.41
DTX2	01/10/00	1520	4,300	6.9	10.1	7.24	0.90
DTX2	04/14/00	1345	4,320	6.9	11.0	7.16	0.76
DTX2	07/07/00	1050	4,510	6.6	12.1	9.16	0.88
DTX2	10/17/00	1005	4,210	6.9	12.5	8.86	0.90
DTX3	01/10/00	1035	1,450	7.0	10.8	8.85	2.7
DTX3	04/11/00	1015	1,840	6.9	9.8	9.17	3.24
DTX3	07/10/00	1030	2,130	6.6	12.4	9.70	3.56
DTX3	10/11/00	1030	2,050	7.0	14.6	10.24	3.24
DTX4	01/06/00	955	3,000	6.6	10.1	7.93	0.90
DTX4	04/11/00	1145	3,070	6.7	8.7	7.25	1.07
DTX4	07/11/00	1010	3,210	6.4	11.6	10.12	1.33
DTX4	10/11/00	1215	3,180	6.7	14.5	12.51	0.96
DTX5	01/06/00	1310	2,950	6.8	10.8	9.78	0.90
DTX5	04/11/00	1345	2,820	6.9	10.3	9.65	0.90
DTX5	07/11/00	1150	3,100	6.6	12.5	10.80	1.23
DTX5	10/16/00	1015	2,950	6.8	12.6	11.97	1.07
DTX6	01/06/00	1530	4,150	6.9	11.7	21.66	0.80
DTX6	04/14/00	1000	4,380	7.0	12.7	21.11	1.01
DTX6	07/13/00	1300	4,500	6.7	13.5	21.90	1.35
DTX6	10/11/00	1400	4,290	7.0	13.4	22.49	1.00
DTX8A	01/13/00	1315	1,850	7.4	11.4	7.97	0.40
DTX8A	04/13/00	1415	1,850	7.3	15.4	7.54	
DTX8A	07/10/00	1505	1,950	7.2	18.8	9.47	0.62
DTX8A	10/16/00	1435	1,830	7.3	15.1	9.58	0.95

Table 12. Field-parameter data associated with ground-water sampling near Deer Trail, Colorado, 2000—Continued [μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter, --, no data because equipment malfunctioned]

Well	Sample date (mm/dd/yy)	Sample time (hhmm)	Specific conduc- tance, field (μS/cm)	pH, field (stan- dard units)	Water temperature (degrees Celsius)	Water level before sampling, depth below measuring point (feet)	Oxygen, dissolved (mg/L)
DTX10A	01/11/00	1130	3,150	7.0	13.2	12.95	0.20
DTX10A	04/17/00	1415	3,120	6.9	17.0	13.03	0.42
DTX10A	07/13/00	1020	3,220	6.8	20.2	13.42	0.26
DTX10A	10/18/00	1040	3,070	7.1	16.2	13.12	3.04
D6	01/11/00	1355	16,000	7.0	12.0	9.35	0.30
D6	04/13/00	945	16,600	6.9	11.7	9.31	0.77
D6	07/11/00	1345	17,500	6.7	13.4	9.44	0.98
D6	10/12/00	1300	16,100	6.9	12.7	9.54	0.70
D13	01/12/00	1005	1,460	7.0	8.5	7.46	0.50
D13	04/17/00	1115	1,390	7.0	9.0	6.59	0.44
D13	07/07/00	1230	1,500	6.8	13.2	8.13	0.76
D13	10/16/00	1215	1,670	7.1	13.8	8.47	1.52
D17	01/07/00	1500	470	7.5	12.1	10.99	1.50
D17	04/10/00	1415	480	7.4	12.3	10.84	2.46
D17	07/06/00	1320	540	7.3	14.1	11.11	1.74
D17	10/10/00	1245	480	7.6	14.4	12.65	0.49
D25	01/12/00	1245	4,300	7.0	11.8	10.32	0.40
D25	04/10/00	1615	4,260	6.9	11.3	10.23	0.66
D25	07/06/00	1500	4,620	6.8	13.2	10.69	0.71
D25	10/10/00	1455	4,910	7.1	13.8	8.81	0.39
D29	01/07/00	1130	4,020	6.7	17.6	154.41	3.0
D29	04/10/00	1215	3,940	6.5	20.0	154.35	3.51
D29	07/06/00	1105	4,090	6.5	23.1	154.35	1.74
D29	10/12/00	1110	3,960	6.6	20.8	154.19	1.08
D30	01/11/00	1600	4,870	6.9	10.7	5.10	0.30
D30	04/13/00	1200	5,060	6.8	10.2	4.75	0.49
D30	07/10/00	1245	5,110	6.7	12.1	6.07	0.59
D30	10/12/00	1450	4,960	7.0	14.4	5.38	0.40

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000

[µS/cm, microsiemens per centimeter at 25 degrees Celsius; mgL, milligrams per liter; µgL, micrograms per liter; per liter; c., less than; ND, no data available; E, value estimated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration candidated for that sample; negative activity reported for some of the nediodremical samples means the sample counts were less than the laboratory background counts for that day]

Chloride, dissolved (mg/L as Cl)	49.7	47.5	49.9	49.1	45.6	40.2	36.0	40.2	17.1	26.0	28.6	28.7	15.2	16.1	17.4	16.1
Acid- neutralizing capacity, titration to pH 4.5, lab (mg/L as CaCO ₃)	319	318	315	316	904	404	405	409	274	314	328	338	353	424	346	348
Sodium (percent)	28	27	28	30	59	28	29	33	15	11	16	17	17	16	18	18
Sodium, dissolved (mg/L as Naj	372	331	355	331	38	360	392	382	58.2	72.8	94.1	5'66	160	158	169	179
Sodium adsorp- tion ratio	4	es	en	4	4	8	4	4	o.	-	-	-	7	7	7	61
Potas- sium, dissolved (mg/Las	3.5	3.3	3.1	3.5	978	7.3	7.8	8.5	6.2	7.3	7.3	7.5	00 00	7.2	7.5	80.00
Magne- sium, dissolved (mg/L as	221	198	209	124	161	351	214	122	6.69	24.2	102	102	6.28	8.08	89.1	95.4
Calcium, dissolved (mg/L as Ca)	477	453	89	467	460	495	472	674	0,1	240	366	263	531	551	526	563
Hardness total (mg/L as CaCO ₃)	2,100	2,000	2,000	1,700	2,000	2,000	2,100	002.1	027	066	1,100	1,100	1,700	1,800	1,700	1,800
pH, labo- ratory (standard units)	7.3	7.2	7.3	7.3	7.1	7.0	7.1	7.0	7.4	7.3	7.4	7.1	7.0	6.9	6.9	6.9
Specific conduc- tance, lab (µS/cm)	4,100	4,150	4,050	4,070	4,300	4,240	4,250	4,230	1,410	1,880	2,020	2,100	2,950	3,140	3,070	3,200
Time (hhmm)	1345	1220	0940	1200	1520	1345	1050	1005	1035	1015	1030	1030	5560	1145	1010	1215
Date (mm/dd/yy)	01/10/00	04/14/00	00/10/100	10/17/00	01/10/00	04/14/00	07/07/00	10/17/00	01/10/00	04/11/00	07/10/00	10/11/00	01/06/00	04/11/00	00/11/20	10/11/00
Well	DTXI	DTX1	DTX1	DTXI	DTX2	DTX2	DTX2	DTX2	DTX3	DTX3	DTX3	DTX3	DTX4	DTX4	DTX4	DTX4

 Table 13.
 Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; μg/L, micrograms per liter; pCi/L, picocuries per liter; c, less than; ND, no data available; E, value estimated by laboratory; --, no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Specific conduc- tance, lab (μS/cm)	pH, labo- ratory (standard units)	Hardness total (mg/L as CaCO ₃)	Calcium, dissolved (mg/L as Ca)	Magne- sium, dissolved (mg/L as Mg)	Potas- sium, dissolved (mg/L as K)	Sodium adsorp- tion ratio	Sodium, dissolved (mg/L as Na)	Sodium (percent)	Acid- neutralizing capacity, titration to pH 4.5, lab (mg/L as CaCO ₃)	Chloride, dissolved (mg/L as Cl)
DTX5	01/06/00	1310	2,900	7.1	1,800	995	86.2	4.6	1	94.6	10	275	8.6
DTX5	04/11/00	1345	2,860	7.1	1,800	268	81.8	4.9	6:	91.0	10	270	7.7
DTX5	07/11/00	1150	2,970	7.1	1,700	559	84.2	4.6	1	120	13	273	10.6
DTX5	10/16/00	1015	2,900	7.1	1,600	523	81.1	5.1	2	148	16	275	10.0
DTX6	01/06/00	1530	4,100	7.2	2,200	468	243	12.8	æ	307	23	270	19.2
DTX6	04/14/00	1000	4,300	7.1	2,200	476	248	13.6	3	309	23	274	21.8
DTX6	07/13/00	1300	4,300	7.3	2,300	468	265	12.6	3	336	24	270	20.6
DTX6	10/11/00	1400	4,290	7.1	2,300	480	263	12.5	3	328	24	270	20.3
DTX8A	01/13/00	1315	1,860	9.7	490	144	32.0	9.9	S	240	51	225	26.9
DTX8A	04/13/00	1415	1,920	7.6	460	135	29.9	7.0	4	219	50	224	27.2
DTX8A	07/10/00	1505	1,870	7.8	510	149	32.4	6.1	5	241	50	216	26.3
DTX8A	10/16/00	1435	1,890	7.5	490	144	31.6	6.2	S	232	50	224	25.6
DTX10A	01/11/00	1130	3,070	7.0	1,800	443	170	112.5	2	150	15	226	17.4
DTX10A	04/17/00	1415	3,180	7.2	1,800	446	171	8.1	2	147	15	223	16.3
DTX10A	07/13/00	1020	3,140	7.2	1,800	433	170	8.8	2	152	16	222	17.9
DTX10A	10/18/00	1040	3,100	7.2	1,800	456	169	8.3	2	153	15	221	16.6

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; μg/L, micrograms per liter; pCi/L, picocuries per liter; https://example.com/readows/liter/, no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Specific conduc- tance, lab (μS/cm)	pH, labo- ratory (standard units)	Hardness total (mg/L as CaCO ₃)	Calcium, dissolved (mg/L as Ca)	Magne- sium, dissolved (mg/L as Mg)	Potas- sium, dissolved (mg/L as K)	Sodium adsorp- tion ratio	Sodium, dissolved (mg/L as Na)	Sodium (percent)	Acid- neutralizing capacity, titration to pH 4.5, lab (mg/L as CaCO ₃)	Chloride, dissolved (mg/L as Cl)
De	01/11/00	1355	16,300	7.2	10,000	400	2,200	11.2	8	1,860	29	889	411
De	04/13/00	0945	16,300	7.2	11,000	446	2,320	12.9	6	2,100	30	640	380
De	07/11/00	1345	16,000	7.2	10,000	414	2,170	12.0	6	2,070	31	889	404
D6	10/12/00	1300	16,200	7.1	10,000	437	2,190	10.4	6	2,110	31	640	419
2	00,017	3001	-	7	730	001	0.73	7	-	0 0 4	ų	0.00	-
	01/17/00	0001	1,410	†. ·	07/	170	, , , , , , , , , , , , , , , , , , ,	† '	٠,	0.00	Ξ ,	0 17	0.1
D13	04/17/00	1115	1,390	7.4	099	179	9.09	2.5	_	57.1	16	253	2.9
D13	00//0//0	1230	1,420	7.7	089	188	52.1	2.8		62.6	17	255	3.2
D13	10/16/00	1215	1,650	7.2	830	226	63.0	2.8	П	2.99	15	256	3.8
D17	01/07/00	1500	492	9.7	210	55.3	18.2	1.6	3.	16.0	41	209	2.3
D17	04/10/00	1415	523	7.7	210	54.9	18.5	1.6	3.	15.4	13	212	2.8
D17	00/90/20	1320	533	7.8	230	56.3	22.5	1.6	4.	15.1	12	215	3.6
D17	10/10/00	1245	484	7.7	220	59.7	18.0	1.6	٠ċ	15.5	13	202	2.9
D25	01/12/00	1245	4,300	7.3	2,500	642	206	7.2	2	260	19	449	108
D25	04/10/00	1615	4,320	7.2	2,400	646	194	7.1	2	240	18	400	119
D25	00/90/20	1500	4,510	7.3	2,600	638	232	7.3	2	286	20	496	8.66
D25	10/10/00	1455	4,900	7.2	2,900	794	219	8.3	3	334	20	585	81.5

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[lµS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; µg/L, micrograms per liter; pCi/L, picocuries per liter; <, less than; ND, no data available; E, value estimated by defaboratory; --, no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Well Date Time number (mm/dd/yy) (hhmm)	Time (hhmm)	Specific conduc- tance, lab (μS/cm)	pH, labo- ratory (standard units)	Hardness total (mg/L as CaCO ₃)	Calcium, dissolved (mg/L as Ca)	Magne- sium, dissolved (mg/L as Mg)	Potas- sium, dissolved (mg/L as K)	Sodium adsorp- tion ratio	Sodium, dissolved (mg/L as Na)	Sodium (percent)	Acid-neutral- izing capacity, titration to pH 4.5, lab (mg/L as CaCO ₃)	Chloride, dissolved (mg/L as CI)
D29	01/2/00	1130	4,020	6.9	2,800	535	346	11.1	1	143	10	293	11.5
D29	04/10/00	1215	4,060	6.7	2,600	521	316	10.9	1	138	10	290	12.3
D29	00/90/20	1105	3,910	8.9	2,600	529	315	10.3	1	137	10	290	13.0
D29	10/12/00	1110	3,980	8.9	2,700	544	313	9.5	-	139	10	283	10.8
D30	01/11/00	1600	4,800	7.1	2,700	428	400	3.9	æ	371	23	364	53.1
D30	04/13/00	1200	4,980	7.2	2,800	425	410	4.0	8	361	22	367	50.0
D30	07/10/00	1245	4,640	7.4	2,600	419	386	3.9	3	369	23	344	47.4
D30	10/12/00	1450	4,870	7.1	2,700	439	389	3.6	3	381	23	346	54.0

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; μg/L, picocuries per liter; less than; ND, no data available; E, value estimated by laboratory; --, no sample submitted; analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Well Date number (mm/dd/yy)	Time (hhmm)	Fluoride, dissolved (mg/L as F)	Silica, dissolved (mg/L as SiO ₂)	Sulfate, dissolved (mg/L as SO ₄)	Nitrogen ammonia plus organic, dissolved (mg/L as N)	Nitrogen ammonia plus organic, total(mg/L as N)	Nitrogen ammonia, dissolved (mg/L as N)	Nitrite plus nitrate (mg/L as N)	Phos- phorus, dissolved (mg/L as P)	Phos- phorus, total (mg/L as P)	Solids, residue on evaporation at 180°C, dissolved (mg/L)	Dissolved solids, sum of constituents (mg/L)
DTX1	01/10/00	1345	0.8	33.9	2,440	0.22	0.23	E.025	2.80	0.073	0.070	4,170	3,810
DTX1	04/14/00	1220	∞.	30.5	2,400	.24	.21	E.017	1.38	.081	.054	4,090	3,670
DTX1	00//0//0	0940	∞.	32.0	2,390	.22	.20	.093	1.29	080	080	4,000	3,700
DTX1	10/17/00	1200	T.	32.2	2,400	E.09	.26	E.047	1.22	.085	080	4,050	3,610
DTX2	01/10/00	1520	٠ċ	15.9	2,430	98.	.62	.549	.05	<.05	<.05	4,200	3,780
DTX2	04/14/00	1345	s.	16.1	2,410	68.	.84	.583	<.04	<.05	<.05	4,160	3,780
DTX2	00/L/0L0	1050	3.	16.6	2,480	88.	.85	.651	<.04	<.05	<.05	4,230	3,870
DTX2	10/17/00	1005	٠ċ:	16.5	2,440	1<.10	86:	.634	<.04	<.05	E.04	4,140	3,740
DTX3	01/10/00	1035	4.	15.7	516	.19	.16	E.014	2.78	<.05	<.05	1,090	1,030
DTX3	04/11/00	1015	4.	16.6	770	<.10	.17	<.029	3.80	<.05	<.05	1,550	1,440
DTX3	07/10/00	1030	4.	10.4	098	.20	.19	<.029	1<.04	<.05	<.05	1,680	1,570
DTX3	10/11/00	1030	4.	17.8	901	.20	.20	E.036	4.23	<.05	>:00	1,760	1,640
DTX4	01/06/00	0955	.2	11.6	1,590	.26	.29	.063	.33	<.05	<.05	2,800	2,620
DTX4	04/11/00	1145	.2	11.5	1,670	<.10	.31	950.	.15	<.05	<.05	3,000	2,760
DTX4	07/11/00	1010	5.	10.7	1,670	.29	.29	.085	4.66	<.05	<.05	2,970	2,720
DTX4	10/11/00	1215	£.	11.6	1,760	.40	.38	.121	6.52	<.05	>:00	3,060	2,880

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; μg/L, picocuries per liter; -, less than; ND, no data available; E, value estimated by laboratory: -., no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Fluoride, dissolved (mg/L as F)	Silica, dissolved (mg/L as SiO ₂)	Sulfate, dissolved (mg/L as SO ₄)	Nitrogen ammonia plus organic, dissolved (mg/L as N)	Nitrogen ammonia plus organic, total (mg/L as N)	Nitrogen ammonia, dissolved (mg/L as N)	Nitrite plus nitrate (mg/Las	Phos- phorus, dissolved (mg/L as P)	Phos- phorus, total (mg/L as	Solids, residue on evaporation at 180°C, dissolved (mg/L)	Dissolved solids, sum of constitu- ents (mg/L)
DTX5	01/06/00	1310	0.2	11.7	1,720	0.12	0.16	0.048	<0.04	<0.05	<0.05	2,920	2,660
DTX5	04/11/00	1345	0.3	11.2	1,670	<0.10	0.18	0.054	<0.04	<0.05	<0.05	2,790	2,600
DTX5	07/11/00	1150	6.	11.2	1,750	.15	.12	.113	<.04	<.05	<.05	2,950	2,710
DTX5	10/16/00	1015	£.	11.9	1,640	.15	.18	.116	<.04	<:05	E.04	2,750	2,590
DTX6	01/06/00	1530	0.4	11.4	2,560	0.12	0.12	0.034	0.26	<0.05	<0.05	4,210	3,790
DTX6	04/14/00	1000	z.	11.2	2,660	11.	14	E.015	.26	<.05	<.05	4,380	3,910
DTX6	07/13/00	1300	4.	11.3	2,730	.10	11.	090.	.23	<.05	<.05	4,440	4,010
DTX6	10/11/00	1400	٠Ċ	11.1	2,680	.14	E.10	.087	.22	<.05	>:00	4,360	3,960
DTX8A	01/13/00	1315	4.	13.0	191	1.6	1.5	1.41	<.04	<.05	<.05	1,420	1,370
DTX8A	04/13/00	1415	4.	12.2	748	1.6	1.5	1.41	<.04	1.12	<.05	1,400	1,320
DTX8A	04/10/00	1505	4.	13.5	757	1.5	1.4	1.39	<.04	<.05	<.05	1,390	1,360
DTX8A	10/16/00	1435	.3	12.9	751	1.3	1.5	1.35	<.04	<.05	>:00	1,410	1,340
DTX10A	01/11/00	1130	0.8	16.6	1,890	4.1	4.1	1.24	<0.0>	<0.05	<0.05	3,100	2,850
DTX10A	04/17/00	1415	6.	16.8	1,880	1.3	1<.10	1.27	<.04	<.05	<.05	3,080	2,830
DTX10A	07/13/00	1020	6.	16.7	1,880	1.4	1.4	1.28	<.04	<.05	<.05	3,090	2,820
DTX10A	10/18/00	1040	∞.	16.8	1,880	1<.10	1.2	1.19	<.04	<:05	90:>	3,070	2,840
D6	01/11/00	1355	L.	20.1	13,000	1.6	1.6	790.	11.7	<.05	E.04	21,000	183,00
9Q	04/13/00	0945	∞.	21.3	12,900	1.5	1.4	.045	13.1	E.04	E.04	20,300	18,600
9Q	07/11/00	1345	∞.	20.6	13,100	1.5	1.4	<.030	14.4	E.03	E.04	21,100	18,700
D6	10/12/00	1300	∞.	8.61	13,100	1.1	1.5	.131	14.6	<.05	E.04	21,500	18,700

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[μS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; μg/L, micrograms per liter; pCi/L, picocuries per liter; <, less than; ND, no data available; E, value estimated by laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Fluoride, dissolved (mg/L as F)	Silica, dissolved (mg/L as SiO ₂)	Sulfate, dissolved (mg/L as SO ₄)	Nitrogen ammonia plus organic, dissolved (mg/L as N)	Nitrogen ammonia plus organic, total (mg/L as N)	Nitrogen ammonia, dissolved (mg/L as N)	Nitrite plus nitrate (mg/L as	Phos- phorus, dissolved (mg/L as	Phos- phorus, total (mg/L as P)	Solids, residue on evaporation at 180°C, dissolved (mg/L)	Dissolved solids, sum of constitu- ents (mg/L)
D13	01/12/00	1005	1.4	12.7	626	.15	.13	.049	<.04	<.05	<.05	1,150	1,110
D13	04/17/00	11115	1.4	11.9	545	.14	.22	<.029	<.04	<.05	<.05	1,080	1,000
D13	00//0//0	1230	1.4	13.6	563	0.14	0.13	0.040	E0.03	<0.05	<0.05	1,110	1,040
D13	10/16/00	1215	1.4	14.1	719	.14	.17	E.027	.05	<.05	<.06	1,340	1,250
D17	01/02/00	1500	1.6	18.2	44.9	E.10	E.10	E.026	2.29	80.	.10	298	294
D17	04/10/00	1415	1.7	17.5	42.4	.12	.12	<.029	3.47	60:	60:	305	298
D17	00/90/20	1320	1.7	18.8	40.6	.16	.17	<.029	5.68	80.	.07	325	315
D17	10/10/00	1245	1.6	19.0	49.3	E.10	E.10	<.049	89:	E.05	.07	287	292
D25	01/12/00	1245	1.0	27.8	2,460	69:0	0.67	0.045	2.2	0.12	0.12	4,520	4,000
D25	04/10/00	1615	1.1	26.3	2,420	.62	.72	.070	2.2	.10	.10	4,390	3,900
D25	00/90/20	1500	1.0	28.5	2,520	.74	<i>LL</i> :	.138	3.0	.15	.15	4,640	4,130
D25	10/10/00	1455	1.0	28.8	2,860	1.11	88.	.110	6.4	.11	.12	5,270	4,710
D29	01/02/00	1130	٠ċ	21.1	2,690	.45	.51	.446	E.02	<.05	E.04	4,260	3,950
D29	04/10/00	1215	9:	20.6	2,630	44.	.47	.428	<.04	<.05	<.05	4,280	3,830
D29	00/90/20	1105	٠ċ	20.4	2,620	.45	.49	.466	E.03	E.04	E.04	4,300	3,830
D29	10/12/00	1110	ĸ	20.2	2,660	.41	.45	.431	<.04	<.05	90.	4,120	3,880
D30	01/11/00	1600	L.	21.8	3,070	.29	.29	.065	<.04	E.04	90.	5,110	4,580
D30	04/13/00	1200	∞.	20.7	3,070	.29	.27	.047	<.04	E.03	E.05	5,070	4,570
D30	07/10/00	1245	∞.	21.0	3,060	.28	.29	.134	<.04	<.05	E.04	5,000	4,520
D30	10/12/00	1450	∞.	21.8	3,100	.30	.34	.120	<.04	<.05	E.06	4,980	4,600

Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued Table 13.

[µS/em_microsiemens per centimeter at 25 degrees Celsius, mg/L, milligrams per liter; µg/L, micrograms per liter; pC/L, piocenties per liter; < less than; ND, no data available; E, value estimated by the laboratory is a combined standard analytical uncertainty; MDC, minimal metertable concentration calculated for that sample; regative activity reported for some of the indiochemical sample means the sample counts were less than the laboratory background courts for that day]

DTX1 04/14/00 1345 7 4 8 4 67 670 620 410 62 8 4 8 4 80 4 80 7 410 620 610 62 8 4 8 4 80 7 410 62 61 9 DTX1 04/14/00 120 4 4 8 4 50 7 410 48 7 11 9 7 11 410 48 4 11 410 410 40 40 50 7 11 9 7 11 9 7 11 9 7 11 9 7 11 9 7 11 11 4 80 4 11 4 80 4 11 4 80 4 11 4 8 4 80 4 11 4 8 4 80 4 11 4 <th>Well</th> <th>Well Date number (mm/dd/yy)</th> <th>Time (hhmm)</th> <th>Aluminum, dissolved (µg/L as Al)</th> <th>Anti- mony, dissolved (ug/L æs Sb)</th> <th>Arsenio, dissolved (μg/L as As)</th> <th>Barium, dissolved (µg/L as Ba)</th> <th>Beryl- lium, dissolved (ug/L as Be)</th> <th>Boron, dissolved (μg/L as B)</th> <th>Bromide, dissolved (mg/L as Br)</th> <th>Cadmium, dissolved (µg/L æs Cd)</th> <th>Chro- mium, dissolved (µg/L as Cr)</th> <th>Cobalt, dissolved (ug/L as Co)</th> <th>Copper, dissolved (µg/L as Cu)</th>	Well	Well Date number (mm/dd/yy)	Time (hhmm)	Aluminum, dissolved (µg/L as Al)	Anti- mony, dissolved (ug/L æs Sb)	Arsenio, dissolved (μg/L as As)	Barium, dissolved (µg/L as Ba)	Beryl- lium, dissolved (ug/L as Be)	Boron, dissolved (μg/L as B)	Bromide, dissolved (mg/L as Br)	Cadmium, dissolved (µg/L æs Cd)	Chro- mium, dissolved (µg/L as Cr)	Cobalt, dissolved (ug/L as Co)	Copper, dissolved (µg/L as Cu)
04/14/00 1220 44 4 8 4 550 33 4.10 4.8 4 07/07/00 0940 4.1 4 8 4 550 33 4.10 4.8 4 10/17/00 120 4 1 8 4 50 33 4.10 4.8 4 01/17/00 1520 4 2 15 4 50 4 5 1 04/14/00 1345 4 4 1 4 5 40 50 4 5 4 1 04/14/00 1345 4 4 1 4 5 4 5 4 5 4 4 6 4 4 5 4 4 5 4 4 5 4 4 4 4 5 4 4 4 4 5 4 4 5 4 4 4 5 4	DTXI	00/01/10	1345	7	a	E	6	Δ	670	0.67	0.2	<1.0	a	ы
900000 610 61 61 8 61 80 61 68 61 68 61 68 61 68 61 68 61 68 61 68 70 61 68 71 61 68 71 61 68 71 61 68 71 61 68 71 61 68 71 61 68 71 61 68 71 61 68 71 61 68 71 <th< td=""><td>DIXI</td><td>04/14/00</td><td>1220</td><td>4></td><td>⊽</td><td>4</td><td>ю</td><td>⊽</td><td>530</td><td>Τ.</td><td><1.0</td><td>N.N</td><td>7</td><td>14</td></th<>	DIXI	04/14/00	1220	4>	⊽	4	ю	⊽	530	Τ.	<1.0	N.N	7	14
10017000 1200 1200 130 20 20 20 20 20 20 20	DTXI	00/20/20	0940	7	⊽	댎	ю	⊽	980	.33	<1.0	×.8	-	6
04/14/00 150 18 4 16 4 310 53 4.0	DTXI	10/17/00	1200	61	a	덦	ю	a	280	0.70	2.0	×.	5	21
04/14/00 1345 c3 c4 E3 c4 290 A0 c4.0 c8 c4 07/07/00 1050 c4 c4 14 c4 350 A1 c4.0 c8 c4 01/07/00 1005 c4	DTX2	01/10/00	1520	2	a	a	16	A	310	R	8	<1.0	80	9
07/07/00 1050 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1 <1	DTX2	04/14/00	1345	۵	⊽	댎	15	7	290	9,	<1.0	×.8	4	Ξ
10,00,0 10,05 2, 2, 2, 2, 2, 2, 2, 2	DTX2	00/20/20	1050	⊽	⊽	E	14	⊽	350	14.	<1.0	v.s	'n	oc
01/10/00 1035 2 4 23 4 210 22 4.10 E6 4 04/11/00 1015 46 4 4 23 4 20 28 4.10 48 4 07/10/00 1030 46 4 21 4 230 29 4.10 4 1 2 20 20 4 2 4 1 1 1 1 1 4 2 20 2 4 2 4 1 1 4 1 4 4 1 4 1 4 4 1 4 4 1 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	DTX2	10/17/00	1005	۵	Ø	回	14	a	98	₩.	0.0	»,	7	=
04/11/00 1015 66 d d d2 23 d d2 29 d d2 6/0 6/8 d d 07/10/00 1030 d d d d d2 21 d d d2 29 d d d d 10/11/00 1030 d d d d d d d d d d d d d d d d d d	DTX3	00/01/10	1035	7	⊽	a	23	⊽	210	13	<1.0	E.6	⊽	М
07/10/00 1030 46 4 4 21 4 230 29 4.10 48 41 10/10/00 1030 4 4 21 4 230 27 4.1 1.7 1 04/10/00 104/10/00 1145 4 4 20 20 22 4.0 4 <td>DTX3</td> <td>04/11/00</td> <td>1015</td> <td>\$</td> <td>7</td> <td>۵</td> <td>23</td> <td>⊽</td> <td>220</td> <td>.28</td> <td><1.0</td> <td>v.S</td> <td>7</td> <td>8</td>	DTX3	04/11/00	1015	\$	7	۵	23	⊽	220	.28	<1.0	v.S	7	8
10/11/00 1030 4 4 4 21 4 230 27 4.1 1.7 1 1 1 1 1 1 1 1 1	Х3	00/01/20	1030	95	⊽	۵	21	⊽	230	29	<1.0	×.8	7	4
04/11/00 0955 <2	DTX3	10/11/00	1030	7	⊽	۵	21	⊽	230	27	7	1.7	-	6
04/11/00 1145 <29 <2 EI 13 <2 270 23 <2.0 <1.0 <2 07/11/00 1010 <1 <2 <2 12 <2 240 29 <2.0 <8 <2 10/11/00 1215 <2 EI <2 14 <1 310 31 EI 2.6 1	X4	00/90/10	0955	۵	a	H	14	a	290	13	2.0	× ×	⊽	v
07/11/00 1010 <1 <2 <2 12 <2 240 29 <2.0 <.8 <2 10/11/00 1215 <2 E.I <2 14 <1 310 .31 E.I 2.6 1	DTX4	04/11/00	1145	8	a	豆	13	a	270	23	0.0	<1.0	۵	9
10/11/00 1215 <2 E.I <2 14 <1 310 .31 E.I 2.6 1	DTX4	00/11/20	1010	⊽	a	۵	12	۵	240	53	0.0	v.S	۵	۲
	X4	10/11/00	1215	۵	E	۵	14	⊽	310	.31	El	2.6	-	9

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[µSkm, microsiemens per centimeter at 25 degrees Celsius, mg/L, milligrams per liter; µg/L, micrograms per liter; pC/L, piccouries per liter; < less than; ND, no data available; E, value estimated by the laboratory; -, no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, miniman detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background courts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Aluminum, dissolved (µg/L as Al)	Anti- mony, dissolved (ug/L as Sb)	Arsenio, dissolved (μg/L as As)	Barium, dissolved (µg/L as Ba)	Beryl- lium, dissolved (ug/L as Be)	Boron, dissolved (μg/L as B)	Bromide, dissolved (mg/L as Br)	Cadmium, dissolved (µg/L as Cd)	Chro- mium, dissolved (µg/L as Cr)	Cobalt, dissolved (ug/L as Co)	Copper, dissolved (µg/L as Cu)
DTX5	00/90/10	1310	2	۵	۵	16	Δ	400	0.20	2.0	6.0	3	9
DTX5	04/11/00	1345	32	A	۵	13	۵	330	91.	0.0	<1.0	m	9
DTX5	00/11/20	1150	7	A	۵	13	۵	390	21	0.0	×.	7	۲
DIXS	10/16/00	1015	۵	-	a	14	⊽	470	25	7	87	61	7
DTX6	00/00/10	1530	8	a	臣	6	A	360	.15	0.0	8,	å	7
DTX6	04/14/00	1000	۵	⊽	7	6	⊽	340	.17	<1.0	οq	7	116
DTX6	07/13/00	1300	m	⊽	۵	6	⊽	340	23	<1.0	3.2	7	6
DTX6	10/11/00	1400	м	∇	۵	ю	⊽	350	71.	E	3.0	-	6
DTX8A	01/13/00	1315	7	∇	a	23	⊽	380	56	<1.0	ES	⊽	61
DTX8A	04/13/00	1415	۵	⊽	۵	21	⊽	240	25	<1.0	×.	7	4
DTX8A	07/10/00	1505	4	⊽	۵	16	⊽	380	.26	<1.0	×.	7	m
DTX8A	10/16/00	1435	8	E03	۵	17	⊽	360	27	7	S.	_	6
DIXIOA	00/11/10	1136	2	7	A	9,	⊽	230	0.25	<1.0	1.5	⊽	e
DTX10A	04/17/00	1415	₹	۵	۵	13	a	240	8	0.0	<1.0	۵	9
DIXIOA	07/13/00	1020	m	7	۵	12	⊽	250	.25	<1.0	3.2	-	9
DIXIOA	10/18/00	1040	å	7	å	12	⊽	250	.28	7	×.	-	v

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[uSGm, microsienens per centimeter at 25 degrees Celsius; mgL, milligrams per liter; µgL, micrograms per liter; pCiL, piocouries per liter; <, less than; ND, no data available; E, value estimated by the laboratory as a corrbined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Aluminum, dissolved (µg/L as Aß	Anti- mony dissolved (µg/L as Sb)	Arsenio, dissolved (ug/L as As)	Barium, dissolved (μg/L as Ba)	Beryl- lium, dissolved (µg/L as Be)	Boron, dissolved (µg/L as B)	Bromide, dissolved (mg/L as Br)	Cadmium, dissolved (ug/L as Cd)	Chro- mium, dissolved (ug/L as Cr)	Cobalt, dissolved (µg/L as Co)	Copper, dissolved (µg/L as Cu)
90	01/11/00	1355	121	⊽	2	7	⊽	068	4.32	<1.0	<4.0	1	51
90	04/13/00	0945	-	7	631	9	7	E750	4.32	<1.0	V.	9	19
90	00/11/20	1345	7	0	H	V	0	820	3.95	0.0	×.8	0	35
8	10/12/00	1300	7	⊽	ю	v,	⊽	970	4.15	Τ.	×.	9	83
DI3	01/12/00	1005	7	⊽	a	8	⊽	80	.18	<1.0	σ,	⊽	6
DI3	04/17/00	1115	0	⊽	a	17	⊽	08	.17	<1.0	×.8	7	5
DI3	00/00/00	1230	7	7	۵	19	7	100	.15	<1.0	E.5	7	2
DI3	10/16/00	1215	7	⊽	a	24	⊽	110	61.	E02	×,	1	m
710	01/02/00	1500	ю	⊽	E	19	⊽	8	80	<1.0	×.	⊽	⊽
D17	04/10/00	1415	7	7	E	58	7	8	8	<1.0	×8	7	7
D17	07/06/00	1320	7	7	E	8	7	06	60	<1.0	×.8	7	-
D17	10/10/00	1245	7	⊽	Ħ	æ	⊽	8	60	7	×.5	-	-
D25	01/12/00	1245	4	⊽	ю	8	⊽	908	1.57	<1.0	2.6	4	6
D25	04/10/00	1615	-	7	ы	19	7	250	1.26	<1.0	0.12	m	Ξ
D25	00/90/20	1500	7	7	ы	17	7	440	50.0°	<1.0	У У	т	10
D25	10/10/00	1455	۵	å	٤	23	å	909	1.38	0.2	4.4	ν,	6

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

JuSkin, microsimens per centimeter at 25 degrees Celsius; mgL, milligrams per liter; µgL, micrograms per liter; pCiL, piccouries per liter; <, less than; ND, no data available; E, value estimated by laboratory; -.. no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MIXC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Well Date Time number (mm/dd/yy) (hhmm)	Time (hhmm)	Aluminum, dissolved (µg/L as All	Anti- mony dissolved (µg/L as Sb)	Arsenio, dissolved (ug/L as As)	Barium, dissolved (μg/L as Ba)	Beryl- lium, dissolved (µg/L as Be)	Boron, dissolved (µg/L as B)	Bromide, dissolved (mg/L as Br)	Cadmium, dissolved (ug/L as Cd)	Chro- mium, dissolved (ug/L as Cr)	Cobalt, dissolved (µg/L as Co)	Copper, dissolved (µg/L as Cu)
D29	00/20/10	1130	9	۵	Δ	6	Δ	180	0.18	<2.0	0.1>	4	9
D29	04/10/00	1215	e	7	H	6	7	190	.18	<1.0	0.10	-	10
D29	07/06/00	1105	7	7	۵	6	7	190	20	<1.0	ьe	7	90
D29	10/12/00	1110	⊽	⊽	۵	90	⊽	300	22	<1.0	×.8	-	7
D30	01/11/00	1600	5	⊽	田	v	⊽	490	.72	<1.0	⊽	-	vı
D30	04/13/00	1200	å	7	М	01	7	440	.70	<1.0	σ;	61	18
D30	07/10/00	1245	<13	7	۵	0	7	450	987	<1.0	у У	cı	12
D30	10/12/00	1450	-	⊽	61	01	⊽	460	छ	<1.0	8.5	61	6

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[µS/cm, microsiemens per certimeter at 25 degrees Celsius; mg/L, miligrams per liter; µg/L, micrograms per liter; pCid., picocunies per liter; <, less than; ND, no data available; E, value estimated by the laboratory as a combined standard analytical uncertainty; MIC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample courts were less than the laboratory background counts for that day]

Zinc, Uranium, dissolved natural (µg/L as (µg/L as Zn) U)	7 54	<14 54	4 48	8 50	7 36	<10 36	4 36	9 34	2 20	<3 28	5 28	2 31	3 32	<11 35	3 27	5 32
Stron- tium, dissolved (µg/L as Sr)	5,960	5,340	5,640	5,460	4,840	5,000	5,450	5,200	2,070	2,770	3,070	2,910	4,180	4,470	4,200	4300
Silver, dissolved (µg/L as Ag)	Q	⊽	⊽	۵	۵	⊽	⊽	A	⊽	7	⊽	7	۵	۵	۵	V
Sele- nium, dissolved (µg/L as Se)	a	ø	E	6	a	4	E	ы	ю	14	14	17	딥	e	E	12
Nickel, dissolved (µg/L as Ni)	22	4	91	31	15	2	15	56	4	9	6	5	⊽	20	10	00
Molyb- denum, dissolved (µg/L as Mo)	9	9	9	7	å	7	61	۵	⊽	7	7	-	۵	۵	۵	-
Mercury, dissolved (µg/L as Hg)	<0.2	<22	<.2	\$	7	<22	7	2	22	ζ2	<.2	ζ2	7	7	<.2	,
Manga- nese, dissoked (µg/L as Mn)	06	77	96	130	3,890	3,700	4,130	3,640	⊽	7	⊽	H	37	33	36	8
Lead, dissolved (µg/L as Pb)	a	⊽	⊽	۵	a	⊽	⊽	۵	⊽	⊽	⊽	∇	å	a	⊽	7
lron, dissolved (μg/L as Fe)	<10	8	8	8	370	400	460	510	<10	<10	8	S	<10	90	8	8
Time (hhmm)	1345	1220	0940	1200	1520	1345	1050	1005	1035	1015	1030	1030	0955	1145	1010	1215
Date (mm/dd/yy)	01/10/00	04/14/00	07/07/00	10/17/00	01/10/00	04/14/00	00/20/20	10/17/00	01/10/00	04/11/00	07/10/00	10/11/00	00/08/00	04/11/00	07/11/00	10/11/00
Well	DIXI	DIXI	DIXI	DIXI	DIX2	DIX2	DIX2	DTX2	DIX3	DIX3	DIX3	DIX3	DIX4	DIX4	DIX4	DYLYA

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

JuSten, microsiemens per certimeter at 25 degrees Celsius; mg/L, milligrams per liter; µg/L, micrograms per liter; pCid., picrocuies per liter; <, less than; ND, no data available; E, value estimated by laboratory; --, no sample submitted; analytical uncertainty is the 2-signa precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	lron, dissolved (µg A. as Fe)	Lead, dissolved (µg/L as Pb)	nese, dissolved (µg/L as Mn)	Mercury, dissolved (µg/L as Hg)	motypedenum, dissolved (µg/L as Mo)	Nickel, dissolved (µg/L as Ni)	nium, dissolved (µg/L as	Silver, dissolved (µg/Las Ag)	Stron- tium, dissolved (μg/Las Sr)	Zinc, dissolved (µg/L as Zn)	Uranium, natural (µg/L as U)
DIXS	00/90/10	1310	100	a	157	<0.2	۵	146	2	å	5,270	3	39
DIXS	04/11/00	1345	130	å	140	22	۵	25	å	۵	5,080	4∆	38
DIXS	07/11/00	1150	8	۵	120	22	۵	10	å	۵	5,110	m	34
DIXS	10/16/00	1015	160	⊽	126	\$	-	6	m	A	4,540	vi	35
DIX6	01/06/00	1530	OI>	å	å	2	Ą	94,	4	A	5,240	vi	37
DIX6	04/14/00	1000	90	7	7	7	7	7	9	7	5,360	₹	38
DIX6	07/13/00	1300	90	7	⊽	22	7	7	Œ	7	5,720	40	4
DIX6	10/11/00	1400	8	⊽	-	2	-	ю	4	۵	5,370	7	38
DIX8A	01/13/00	1315	8	⊽	180	2	⊽	9	å	7	2,460	2	⊽
DIXSA	04/13/00	1415	130	7	181	2	7	7	Q	7	2,320	۵	7
DIXSA	07/10/00	1505	071	⊽	181	<.2	7	e	Q	7	2,520	₹	7
DIXSA	10/16/00	1435	220	⊽	172	\$	_	5	۵	7	2,420	7	7
DIXIOA	00/11/10	1130	3,850	⊽	1158	40.2	⊽	6	Q	⊽	5,450	5	⊽
DIXIOA	04/17/00	1415	3,970	å	327	\$	۵	121	۵	۵	5,430	<12	۵
DIXIOA	07/13/00	1020	4,130	7	290	<2	-	m	۵	7	5,490	4	7
DIXIOA	10/18/00	1040	4,110	⊽	394	<.2	-	-	a	۵	5,510	4	M

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[pS/cm, microsiemens per centimeter at 25 degrees Celsius, mg/L, millignams per liter, pg/L, micrograms per liter; pCi/L, picocuries per liter; <, less than; ND, no data available; E, value estimated by laboratory; --, no sample submitted; analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample courts were less than the laboratory background counts for that day]

Pb)	dissolved dissolved (μg/Las (μg/Las Fe) Pb)
3,970	3,97(
157.1	157.
3,680	3,68
3,760	3,76
28	
82	~
76	1>
48	4
249	1 249
213	<1 21:
298	≥ 38
347	5
3,100	3,10
3,120	4 3,13
3,290	<1 3,29
3.790	

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[µS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, millignams per liter; µg/L, micrograms per liter; pCi/L, picocuries per liter; c, less than; ND, no data available; E, value estimated by the laboratory; -.. no sample submitted; analytical uncertainty is the 2-signa precision estimate calculated by the laboratory as a combined structured analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample courts were less than the laboratory background counts for that day]

Well	Well Date number (mm/dd/yy)	Time (hhmm)	Iron, dissolved (µg/L as Fe)	Lead, dissolved (µg/Las Pb)	Manga- nese, dissolved (ug/L as Mn)	Meroury, dissolved (ug/L as Hg)	Molyb- denum, dissolved (µg/L as Mo)	Nickel, dissolved (µg/L as Ni)	Sele- nium, dissolved (µg/L as Se)	Silver, dissolved (ug/L as Ag)	Stron- tium, dissolved (μg/L as Sr)	Zino, dissolved (µg/L as Zn)	Uranium, natural (ug/L as U)
D29	00/20/10	1130	7,210	۵	817	<2	۵	1>1	۵	4	5,830	œ	۵
D29	04/10/00	1215	5,970	7	813	7	-	7	EI	۵	5,840	19	7
D29	00/90/20	1105	7,080	7	788	\$	⊽	24	H	7	5,810	6	7
D29	10/12/00	1110	01979	⊽	88	7	-	7	6	7	5,740	13	⊽
D30	00/11/10	1600	Я	⊽	149	2	2	15	a	⊽	5,880	4	17
D30	04/13/00	1200	E40	7	293	7	en	5	'n	7	010'9	>16	40
D30	07/10/00	1245	100	7	253	\$	en	14	۵	7	5,740	11>	35
D30	10/12/00	1450	1560	7	380	\$	2	4	e	⊽	5,780	9	35

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

Well	Date (mm/dd/yy)	Time (hhmm)	Plutonium-238, whole water (pGi/L)	Analytical uncertainty, Plutonium-238 (pCi/L)	MDC, Plutonium-238 (pGi/L)	Plutonium- 239+240, whole water (pGiA)	Analytical uncertainty, Plutonium-239+240, (pCi/L)	MDC, Plutonium- 239+240 (pCi/L)
DIXI	01/10/00	1345	0	0.012	0.013	-0.002	0.004	0.027
DIXI	04/14/00	1220	1	1	1	1	1	;
DIXI	00/20/20	0540	.1	1		1	1	1
DTXI	10/17/00	1200	0	0.012	0.014	0.005	010'0	0.014
DTX2	01/10/00	1520	800'0	0.016	0.022	0	02070	0.022
DTX2	04/14/00	1345	1	1	1	1	:	1
DTX2	00/20/20	1050	te	r			F	;
DTX2	10/17/00	1005	0	0.007	0.008	0	2000	0.008
DTX3	01/10/00	1035	0	0.004	0.004	0,001	0.004	0.009
DIX3	04/11/00	1015	1	1	;	1	1	
DIX3	07/10/00	1030	1	1	1	1	1	
DIX3	10/11/00	1030	0	0.008	0.009	0.002	2000	0.018
DTX4	00/90/10	9955	-0.003	80000	0.027	0.006	8000	0.008
DTX4	04/11/00	1145	ŀ	18	1	1	1	
DTX4	07/11/00	1010	1	ा	1	1	1	1
DIXA	10/11/00	1215	-0.002	0.003	0.024	0000	0.012	0.012

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[u8/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; pg/L, micrograms per liter; pC/L, piccouries per liter; < less than; ND, no data available; E, value estimated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Plutonium-238, whole water (pGi/L)	Analytical uncertainty, Plutonium-238 (pCi/L)	MDC, Plutonium-238 (pCi/L)	Plutonium- 239+240, whole water (pCi/L)	Analytical uncertainty, Plutonium-239+240, (pCi/L)	MDC, Plutonium- 239+240 (pCi/L)
DTX5	00/90/10	1310	0	0.013	0.014	0.003	0.012	0.030
DTX5	04/11/00	1345	1	ı	1	1	1	
DTX5	07/11/00	1150	1	1	;	1	1	1
DIXS	10/16/00	1015	-0.001	0.002	0.016	0.003	9000	0.008
DTX6	00/90/10	1530	-0.003	9000	0.042	0	810.0	0.020
DTX6	04/14/00	1000	1	E	1	1	1	,
DTX6	07/13/00	1300	1	1	ा	1	1	1
DTX6	10/11/00	1400	0	0.004	0.004	0.002	0000	0.004
DTX8A	01/13/00	1315	0	9000	9000	-0.002	0000	0.017
DIXSA	04/13/00	1415	+	1	1	1	1	1
DIXSA	07/10/00	1505	ı	1	1	1	1	1
DTX8A	10/16/00	1435	0	0.004	0.005	-0.001	100'0	0.010
DIXIOA	00/11/10	1130	0.003	0.012	0.030	0.003	0.012	0.030
DIXIOA	04/17/00	1415	,	,		1	1	1
DTX10A	07/13/00	1020	1	ī	1	1	1	
4 01 2010								

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[JuNem, microsiemens per centimeter at 25 degrees Celsius, mgl.L. milligrams per liter; pg/L. micrograms per liter; pCVL, picocuries per liter; < less than; ND, no data available; E, value estimated by the laboratory as a combined standard malytical uncertainty; MDC, minimum detectable concentration calculated for that sample, regative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Date (mm/dd/yy)	Time (hhmm)	Plutonium-238, whole water (pCi/L)	Analytical uncertainty, Plutonium-238 (pCi/L)	MDC, Plutonium-238 (pCi/L)	Plutonium- 239+240, whole water (pCi/L)	Analytical un certainty, Plutonium- 239+240, (pCi/L)	MDC, Plutonium- 239+240 (pCi/L)
D6	00/11/10	1355	9000	0.020	0.051	-0.004	0.007	190'0
D6	04/13/00	0945		,	ı		1	
9G	07/11/00	1345	1		1		:	
90	10/12/00	1300	0	0.007	0.008	1000	0.007	0.020
D13	01/12/00	1005	100'0-	0.002	0.013	0000	0.004	0.013
D13	04/17/00	1115	1	1	r	1	1	1
D13	00/00/00	1230	1	1	1	1	1	1
D13	10/16/00	1215	100'0	0.009	0.026	0.002	0.008	0.022
71G	00/20/10	1500	100'0-	0.001	0.008	0	0.004	1000
D17	04/10/00	1415			1	1	1	
D17	07/06/00	1320	1	1	1		1	
D17	10/19/00	1245	100'0-	0.002	0.011	0.002	0.003	100'0
D25	01/12/00	1245	-0.004	9000	0.034	100'0	0.011	150'0
D25	04/10/00	1615	1	;	1		:	1
D25	07/06/00	1500		1	1		1	1
D25	10/10/00	1455	0	0.008	0000	0,003	0000	0000

Table 13. Water-quality data for monitoring wells near Deer Trail, Colorado, 2000—Continued

[JuS/cm, microsienzens per centimeter at 25 degrees Celsius, myL., milligrams per liter; µg/L, micrograms per liter; pC/L, piccouries per liter; < less than; ND, no data available; E, value estimated by becratory. -., no sample submitted, analytical uncertainty is the 2-signra precision estimate calculated by the laboratory as a combined standard analytical uncertainty; MDC, minimum detectable concentration calculated for that sample, regative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Well	Well Date Time number (mm/dd/yy) (hhmm)	Time (hhmm)	Plutonium-238, whole water (pGi/L)	Analytical uncertainty, Plutonium-238 (pGiA.)	MDC, Plutonium-238 (pGi/L)	Plutonium- 239+240, whole water (pCi/L)	Plutonium- 239+240, (pCi/L)	
D29	00/20/10	1130	-0.002	0.005	0.034	9000	0.012	
D29	04/10/00	1215	ı	1	1	1	1	
D29	00/00/00	1105	1	;	1	1	1	
D29	10/12/00	01110	010'0-	0.010	0.051	-0.002	0.005	
D30	00/11/10	091	0	0.019	0.021	800'0	0.016	
D30	04/13/00	1200	10	ı	18	ı	1	
D30	00/10/00	1245	1	,	1		1	1
D30	10/12/00	1450	-0.002	0.004	0.031	-0.002	0.004	0.031

1 Value is significantly different from historical or subsequent data at the same site, and analytical bias is sispecied. However, issufficient evidence from laboratory to reject or change value.

Table 14. Quality-control data for blank samples associated with ground-water samples collected near Deer Trail, Colorado, 2000

[µS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter, µg/L, micrograms per liter; pCi/L, picocuries per liter, Q, equipment blank; E, field blank; S, source-water blank; c, less than; E, value estimated by laboratory, --, no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined structural uncertainty: MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Chloride, dissolved (mg/L as Cl)	<0.3	\$:	Ş	Ş	Ş	:	Ş	:		77	۲.
Acid-neutral- izing capacity, titration to pH 4.5 lab (mg/L as CaCO ₃)	2	2	1	2	2	2	,	7	1		2	1
Sodium, dissolved (mg/L as Na)	40.1	7.7	7	7	77	7	7	7	77	7	7	7
Potas- sium, dissolved (mg/L as K)	<0.2	2	1	\$	\$	\$;	\$	1	1	\$	\$
Magnesium, dissolved (mg/L as Mg)	<0.01	<.01	<.01	<.01	<.01	20.	10.	<.01	<.01	<01	×01	<.01
Calcium, dissolved (mg/L as Ca)	<0.02	<.02	<.01	<.02	<.02	80.	70.	<.02	<.01	<.01	<.01	<.01
pH, labora- tory (stan- dard units)	7.8	7.3	1	7.4	80	7.5	1	8.4	1	1	7.9	8.1
Specific conduc- tance, lab (µS/cm)	4	Ξ	1	4	E	E	ŀ	6	1	1	4	4
Well	D17	D6	9Q	DTX5	D13	DIXIOA	DIXIOA	DTX6	DTX6	None	DTX3	DTXI
Time (hhmm)	1430	1400	1405	1315	1045	1015	1016	1240	1045	1050	0940	1140
Date (mm/dd/yy)	00/20/10	01/11/00	01/11/00	04/11/00	04/17/00	04/21/00	04/21/00	07/13/00	07/14/00	07/14/00	10/11/00	10/17/00
Blank type	ц	щ	$\mathbf{F}_{\mathbf{l}}$	L	Ľ,	o	70'	Ľ,	s_1	N_1	Ĭ.	L

Quality-control data for blank samples associated with ground-water samples collected near Deer Trail, Colorado, 2000—Continued Table 14.

[JSkm, microsiemens per centimeter at 25 degrees Celsius; mgL, milligrams per liter; µg/L, micrograms per liter; µG/L, picocuries per liter; Q, equipment blank; F, field blank; S, source-water blank; <, less than; E, value estimated by laboratory; -, no sample submitted; analytical uncertainty is the 2-sigma precision estimate by the laboratory as a combined standard analytical incertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the Informatory background counts for that day]

Blank type	Date Fluoride, (mm/dd/yy) (mg/L æ F)	Fluoride, dissolved (mg/L as F)	Silica, dissolved (mg/L as SiO ₂)	Sulfate, dissolved (mg/L as SO ₄)	Nitrogen ammonia plus organie, dissolved (mg/L as N)	Nitrogen ammonia plus organio, total (mg/L as N)	Nitrogen ammonia, dissolved (mg/L as N)	Nitrite plus nitrate (mg/L as N)	Phosphorus, dissolved(mg/L as P)	Phosphorus, total (mg/L as P)
н	01/07/00	<0.1	<0.1	<0.3	E10	<0.10	E.021	<0.04	<0.05	<0.05
ш,	01/11/00	8.1	9.1	<0.3	<0.10	<.10	<0.029	E0.02	<.05	<.05
74	01/11/00	;	7	1	1	1			1	ı
щ	04/11/00	7	7	<3	<.10	<.10	<.029	90	<.05	\$0.5
щ	04/17/00	7	7	<.3	<.10	<.12	<.029	<04	<.05	<.05
0	04/21/00	7	E.04	5	<.10	<.10	<.029	<.04	<.05	<.05
·0	04/21/00	;	7		1	ı	1	,	1	ı
щ	07/13/00	7	7	<3	<.10	<.10	<.029	<.04	<.05	<.05
5,1	07/14/00	;	77		1	1	1	:	1	1
5.	07/14/00	;	7	1	1	ı	1	:	1	ı
щ	10/11/00	7	7	Е1	<.10	<.10	<.049	<.04	<.05	>:00
щ	10/17/00	2	7	5	<.10	<.10	610.>	<.04	<.05	>:00

Table 14. Quality-control data for blank samples associated with ground-water samples collected near Deer Trail, Colorado, 2000—Continued

Less tran; E, value estimated by Informatory; —, no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the Informatory as a corriend standard analytical uncertainty; MEC, minimum detectable concentration calculated for that sample; negative activity reported for some of the indiscremental samples means the sample counts were less than the laboratory background. JuSkin, microsiemens per centimeter at 25 degrees Celsius; mgL, milligrams per liter; µgL, micrograms per liter; pCVL, pieceuries per liter; Q, equipment blank; E, field blank; S, source-water blank; counts for that day]

Blank type	Date (mm/dd/yy)	Chromium, dissolved (µg/L as Cr)	Cobalt, dissolved (µg/L as Co)	Copper, dissolved (µg/L as Cu)	lron, dissolved (ug/L as Fe)	Lead, dissolved (µg/L as Pb)	Manganese, dissolved (μg/L as Mn)	Mercury, dissolved (µg/L as Hg)	Molybdenum, dissolved (ug/L as Mo)	Nickel, dissolved (µg/L as Ni)	Selenium, dissolved (µg/L as Se)
ы	00/20/10	8.0>	⊽	⊽	<10	<1	- 1	<0.2	1>	1>	۵
щ	00/11/10	×.8	⊽	⊽	<10	7	⊽	<.2	7	7	å
-j.,	01/11/00	7	⊽	⊽	Ø	7	7	;	7	7	;
ш,	04/11/00	8.>	⊽	⊽	OI>	7	⊽	ζ,	7	7	a
щ	04/17/00	8.>	⊽	⊽	OI>	7	7	7	7	7	a
0	04/21/00	×.8	⊽	16	E10	7	⊽	2	7	⊽	۵
-o	04/21/00	42	⊽	16	01	⊽	⊽	:	7	⊽	10
щ	07/13/00	E.6	⊽	⊽	OI>	7	⊽	2	7	7	a
s_1	07/14/00	<2	⊽	⊽	۵	7	⊽	;	7	7	1
s_1	07/14/00	2	⊽	⊽	۵	~	⊽		7	7	1
щ	10/11/00	×.8	⊽	⊽	OI>	7	⊽	7	7	7	å
ш,	10/17/00	×.8	⊽	E.22	OI>	7	7	\$	7	7	a

Quality-control data for blank samples associated with ground-water samples collected near Deer Trail, Colorado, 2000—Continued Table 14.

[µS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; µg/L, micrograms per liter; pCi/L, picocaries per liter; Q, equipment blank; E, field blank; S, source-water blank; <, less than; E, value estimated by laboratory; -, no sample submitted; analytical uncertainty is the 2-sigma precision estimate by the laboratory as a combined standard analytical uncertainty, MDC, minimum detectable concentration calculated for that sample, negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Blank type	Date (mm/dd/yy)	Solids, residue on evaporation at 180°C, dissolved (mg/L)	Aluminum, dissolved (µg/L as Al)	Antimony, dissolved (μg/L as Sb)	Arsenic, dissolved (ug/L as As)	Barium, dissolved (µg/L as Ba)	Beryllium, dissolved (µg/L as Be)	Boron, dissolved (ug/L as B)	Bromide, dissolved (mg/L as Br)	Cadmium, dissolved (ug/L as Cd)
IL.	00/20/10	<10	22	⊽	<2.0	⊽	⊽	05	<0.01	<1.0
Ŀ	00/11/10	<10	⊽	⊽	2.0	⊽	⊽	20	<.01	<1.0
TL.	01/11/00	1	⊽	⊽	ı	⊽	⊽	Q	1	ζ3
ш	04/11/00	<10	€	7	0.0	⊽	⊽	20	<.01	<1.0
ы	04/17/00	<10	7	7	2.0	⊽	⊽	20	<.01	<1.0
0	04/21/00	<10	7	7	2.0	⊽	⊽	20	<.01	<1.0
50	04/21/00	1	7	7	10	⊽	⊽	۵	1	5
ь	07/13/00	<10	7	7	0.0	7	7	20	<.01	<1.0
N ₂	07/14/00	1	⊽	7	1	⊽	⊽	۵	,	ζ3
N ₁	07/14/00	1	⊽	7	1	⊽	⊽	۵	1	\$
Ľ.	10/11/00	<10	7	7	2.0	7	~	010	<.01	7
Ľ,	10/17/00	<10	7	E03	2.0	7	⊽	OT>	<.01	7

Quality-control data for blank samples associated with ground-water samples collected near Deer Trail, Colorado, 2000—Continued Table 14.

[JS/cm, microsiemens per centimeter at 25 degrees Celsius; mg/L, milligrams per liter; pCi/L, piccounies per liter; Q, equipment blank; F, field blank; S, source-water blank; c, less than; E, value estimated by laboratory; -., no sample submitted; analytical uncertainty is the 2-sigma precision estimate calculated by the laboratory as a combined stracked analytical uncertainty; MDC, minimum detectable concentration calculated for that sample; negative activity reported for some of the radiochemical samples means the sample counts were less than the laboratory background counts for that day]

Blank type	Date (mm/dd/ yy)	Silver, dissolved (µg/L as Ag)	Stron- tium, dissolved (µg/L as Sr)	Thallium, dissolved (µg/L)	Zino, dissolved (µg/L as Zn)	Uranium, natural (ug/L as U)	Plutonium- 238 (pCVL)	Analytical uncer- tainty, Plutonium- 238 (pCi/L)	MDC, Pluto- nium-238 (pCi/L)	Plutonium- 239+ 240 (pCi.L)	Analytical uncertainty, Plutonium- 239+240 (pCi/L)	MDC, Plutonium- 239+240 (pCi/L)
F	00/20/10	7	<1.0	;	7	7	0	0000	0.004	1000	0.003	0000
Ŀ,	01/11/00	7	<1.0	;	⊽	7	-0.001	0.001	0.008	0	0.004	0,004
74	01/11/00	7	7	77	7	⊽	1				1	:
14	04/11/00	7	<1.0	1	⊽	⊽	:	1	:	:	18	:
Ŀ,	04/17/00	7	<1.0	1	27	⊽	:	1	:		ा	;
0	04/21/00	7	ES	1	Q	7	20,012	0.008	0.009	100'0	0.002	0000
5	04/21/00	7	DQ.	7	-	⊽	1	1			ı	
14	07/13/00	7	<1.0	1	⊽	⊽			:	1	1	:
N.	07/14/00	7	J	7	⊽	⊽	1	1	:	1	1	:
N,	07/14/00	7	7	7	7	7	1				1	:
H	10/11/00	7	× 8	;	7	7	0000	0.004	0.004	0,002	0.005	60000
ы	10/17/00	7	8 >	;	⊽	7	1000	0.003	0.009	0	0.004	0000

Blank sample for low-level trace-element analysis.

Analytical bias is suspected. However, insufficient evidence from laboratory to reject or charge value.

[RPD, relative percent difference, which is defined as [(sample value-replicate value)/((sample value + replicate value)/((sample valu Table 15. Comparison of water-quality data for replicate and regular ground-water samples collected near Deer Trail, Colorado, 2000

	Well number		DIX			D29			DTX3	
	Date		01/06/00			00/00/10			01/10/00	
	Time	9960	1000		1130	1135		1035	1040	
Property or constituent		Sample	Replicate	RPD	Sample	Replicate	RPD	Sample	Replicate	RPD
Specific conductance, lab (µS/cm at 25 °C)		2,950	2960	0	4,020	4,010	0	1,410	1,410	0
pH, WH, Jabotzatory (standard units)		7.0	7.0	0	6.9	6.8	-	7.4	7.4	0
Hardness total (mg/L as CaCO ₃)		1,700	1,700	0	2,800	2,700	4	720	099	6
Calcium, dissolved (mg/L as Ca)		531	523	7	535	526	61	07.1	158	-
Magnesium, dissolved (mg/L as Mg)		869	84.0	6	346	327	9	6.69	63.6	6
Sodium, dissolved (mg/L as Na)		160	158	1	143	140	7	58.2	54.0	90
Potassium, dissolved (mg/L as K)		90 90	6.8	7	11.1	11.6	7	6.2	6.0	6
Acid neutralizing capacity, titration to 4.5, lab (mg/L as CaCO ₃)	(mg/L as CaCO ₃)	353	393	-11	293	292	0	274	274	0
Sulfate, dissolved (mg/L as SO4)		1,590	1,630	-5	2,690	2,710	7	516	516.86	0
Chloride, dissolved (mg/L as Cl)		15.2	15.6	ņ	11.5	11.6	7	17.1	17.2	7
Fluoride, dissolved (mg/L as F)		7	7	9	v,	4.	13	4.	₹.	0
Brotmide, dissolved (mg/L as Br)		22	22	0	.18	7	ņ	13	13	0
Silica, dissolved (mg/L as SiO ₂)		911	11.8	-1	21.1	20.8	-	15.7	15.4	2
Solids, residue on evaporation at 180°C, dissolved (mg/L)	ved (mg/L)	2,800	2,850	-2	4,260	4,270	0	1,090	1,090	0
Dissolved solids, sum of constituents (mg/L)		2,620	2,670	-	3,950	3,930	-	1,030	010'1	5
Nitrite plus nitrate (mg/L as N)		33	.32	3	E.02	E.02	0	2.78	2.78	0
Nitrogen ammonia, dissolved (mg/L as N)		590	60.	-31	9446	947	0	E014	<.029	R
Nitrogen ammonia plus organic, total (mg/L as N)	(X s	29	.30	-5	51	8	7	.16	.14	10
Nitrogen ammonia plus organic, dissolved (mg/L as N)	yL as N)	26	.25	9	45	84.	9-	.19	.14	32
Phosphorus, total (mg/L as P)		\$0°>	<.05	0	E.O.	<05	ND	<.05	<05	0
Phoenhorne discolard (mail no D)		20.00	2000	0	20.00	ECOLO	CIN	2000	20.00	0

[RPD, relative percent difference, which is defined as [sample value-replicate value*/(sample value + replicate value*/2] × 100; relative percent difference was calculated on uncounded data; -, not analyzed; ND, not determined because data were less than the minimum reporting limit or estimated; c, less than; E, value estimated by laboratory] Table 15. Comparison of water-quality data for replicate and regular ground-water samples collected near Deer Trail, Colorado, 2000—Continued

	Well number		DTX4			D29			DTX3	
	Date		01/06/00			01/02/00			01/10/00	
	Time	9960	1000		1130	1135		1035	1040	
Property or constituent		Sample	Replicate	RPD	Sample	Replicate	RPD	Sample	Replicate	RPD
Aluminum, dissolved (µg/L as Al)		۵	42	0	9	5	18	2	1>	Q
Antimony, dissolved (µg/L as So)		۵	۵	0	۵	۵	0	7	7	0
Absenic, dissolved (µg/L as As)		El	۵	Q.	å	۵	0	۵	۵	0
Barium, dissolved (µg/L as Ba)		14	4	0	6	6	0	23	23	0
Beryllium, dissolved (µg/L as Be)		å	۵	0	å	۵	0	7	7	0
Boron, dissolved (Itg/L as B)		290	280	6	180	190	4	210	210	0
Cadmium, dissolved (µg/L as Cd)		0.0	200	0	2.0	<2.0	0	7	7	0
Chromium, dissolved (µg/L as Cr)		×.×	E.7	Q.	<1.0	<1.0	0	E.6	»	g
Cobalt, dissolved (µg/L as Co)		7	⊽	0	۵	۵	0	7	⊽	0
Copper, dissolved (µg/L as Cu)		5	5	0	9	7	-15	М	7	9
Iron, dissolved (µg/L as Fe)		<10	Ð	Q	7,210	7,110	-	<10	<10	0
Lead, dissolved (Ug/L as Pb)		۵	۵	0	۵	۵	0	7	⊽	0
Manganese, dissolved (µg/L as Mn)		3.7	38	3	817	818	0	7	7	0
Mercuty, dissolved (µg/L as Hg)		<.2	<.2	0	<.2	<2	0	<2	<2	0
Molybdenum, dissolved (µg/L as Mo)		۵	۵	0	۵	♡	0	7	⊽	0
Nickel, dissolved (µg/L as Ni)		7	⊽	0	<u>-</u>	<u>y</u>	0	4	4	0
Selenium, dissolved (µg/L as Se)		Œ	E	0	۵	۵	0	D0	90	0
Silver, dissolved (µg/L as Ag)		۵	۵	0	۵	۵	0	7	⊽	0
Stronium, dissolved (µg/L as Sr)		4,180	4,120	-	5,830	5,760	-	2,070	2,010	e
Zinc, dissolved (µg/L as Zn)		e	60	0	ю	ю	0	7	5	0
Uranium, natural (µg/L as U)		32	32	0	۵	\$	0	30	8	0
Plutonium-238, whose water (pG/L as Pa)	Pu)	-0.003	0	200	0.002	0	200	0	0.001	-200
Plutonium-239-240, whole water (pGAL as Pu)	Lus Pu)	0.000	0	200	9000	c	2000	1000	10009	N

[RPD, relative percent difference, which is defined as [(sample value-replicate value+ replicate value+ replicate value) x 100; relative percent difference was calculated on unrounded data; --, not analyzed; ND, not determined because data were less than the minimum reporting limit or estimated; <, less than; E, value estimated by laboratory] Table 15. Comparison of water-quality data for replicate and regular ground-water samples collected near Deer Trail, Colorado, 2000—Continued

9945 RPD Sample 0 16,300 1 3 11,000 1 3 446 3 2,320 1 2,100 4 12,9 0 640 1 12,900 1 4 380 0 4,32 4 21,3 1 20,300 2 1 20,300 1 0 13,1 0 015	DTX3	90			D25	
Time folia folia 6845 Sample Replicate RPD Sample 1,880 1,880 0 16,300 1 7,3 7,3 7,3 1,000 1 72,8 72,3 1,100 1 72,8 72,3 1,100 1 72,8 72,3 1,100 1 73, 7,0 4 1,2,00 1 25,0 25,1 4 380 1 25,0 25,1 4 380 1 25,0 25,1 4 380 1 25,0 1,50 1,50 1 25,0 1,50 1,410 2 1,0,30 1 25,0 2,0 1,50 1,410 2 1,410 1 25,0 2,0 2,0 1,51 1 25,0 2,0 1,51 1 25,0 3,80 0 1 25,0	04/11/00	04/13/00	ĺ		00/90/20	
Sample Replicate RPD Sample 1,880 1,880 0 16,300 1 7.3 7.3 0 7.2 1 240 232 3 1400 1 728 723 1 1000 1 728 723 1 1 2,100 1 728 729 1 1 2,100 1 250 25.1 4 129 1 250 25.1 4 380 1 250 25.1 4 380 1 250 25.1 4 380 1 250 25.1 4 380 1 250 25.1 1 20,300 1 250 1,500 1,410 2 1,410 2 1,410 1 250 2020 0 13.1 1 250 2020 0 13.1 1 250 2020 0 13.1 1 250 2020 0 13.1 1 250 2020 0 13.1 1 250 2020 0 15.0 1 250 2020 0 15.0 1 250 2020 0 15.0 1 250 2020 0 1.5 1 250 2020 0 1		0980		1500	1505	
1,880 1,880 0 16,300 7.3 7.3 0 7.2 990 990 3 11,000 240 232 3 446 72.8 72.3 1 2,100 72.8 72.3 1 2,100 73.14 314 0 640 70.0 762 1 12,900 25.0 25.1 4 380 4 4 0 8 25.0 25.1 4 380 15.0 15.9 1 12,900 25.0 15.0 1 12,900 25.0 15.0 1 12,900 25.0 15.0 1 12,900 25.0 15.0 1 12,900 25.0 15.0 1 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25.0 15.0 12,900 25	RPD	e Replicate	RPD	Sample	Replicate	RPD
its) 7.3 7.3 0 7.2 990 960 3 11,000 240 232 3 446 Mg) 728 72.3 1 2,100 7.2 7.3 7.0 4 12,90 20 to 4.5, lab (mg/L as CaCO ₃) 314 314 0 640 250 25.1 4 380 250 25.1 4 380 180°C, dissolved (mg/L) 1,550 1,410 20,300 cnts (mg/L) 1,550 1,530 1 1,500 3.80 4.32 indicabled (mg/L as N) 1,71 1,410 20,300 cuts (mg/L as N) 4,00 1,410 2 1,8500 3.80 4.32 indicabled (mg/L as N) 4,00 1,510 dissolved (mg/L as N) 4,00 4,00 1,51 con to dissolved (mg/L as N) 4,00 4,00 1,51 con to dissolved (mg/L as N) 4,00 4,00 1,51	0	16,300	0	4,510	4,490	0
990 960 3 11,000 240 232 3 446 Mg) 942 91,11 3 2,320 72.8 72.3 1 2,100 73 7.0 4 12,9 co to 4.5, lab (mg/L.as Ca.Co.3) 314 314 0 640 770 762 1 12,900 780 25.1 4 380 780 25.1 4 380 780 25.1 4 380 780 25.1 1,410 20,300 cuts (mg/L.s N) 1,410 1,410 2 18,600 dissolved (mg/L.s N) 7.0 4 1,410 780 25.1 4 380 780 25.1 4 380 780 25.1 4 380 780 25.1 4 380 780 25.1 4 380 780 25.1 4 380 780 20.300 780 20.300 780 20.300 780 20.300 780 20.300 780 20.300 780 20.300 780 20.300 780 20.300 780 20.300	0	7.2	0	7.3	7.3	0
Mg) 94.2 91.1 3 2.320 72.8 72.3 1 2.320 72.8 72.3 1 2.320 72.8 72.3 1 2.300 73 770 762 1 12.900 750 25.1 4 380 750 25.1 4 380 750 15.90 15.90 cuts (mg/L.ss N) 1,440 1,410 2 18,600 total (mg/L.ss N) 217 < 20.9 dissolved (mg/L.ss N) 218 dissolved (mg/L.ss N	В	10,000	10	2,600	2,500	4
Mg) 728 728 729 729 729 729 729 729 720 720 720 720 720 720 720 720 720 720	e	431	m	859	624	61
72.8 72.3 1 2,100 73 7.0 4 12.9 co to 4.5, lab (mg/L as CaCO ₂) 314 314 0 640 260 25.1 4 380 28 28 0 4.32 18.0°C, dissolved (mg/L) 1,550 1,530 1,610 18.0°C, dissolved (mg/L) 1,440 1,410 2 20,300 conts (mg/L) 1,440 1,410 2 18.600 dissolved (mg/L as N) 1,7 1,40 1,410 2 18.600 dissolved (mg/L as N) 1,7 1,40 1,410 2 18.600 dissolved (mg/L as N) 2,02 2 1,530 1,530 cost (mg/L as N) 2,02 2 1,530 1,540 dissolved (mg/L as N) 2,17 1,40 1,410 cost (mg/L as N) 2,17 1,410 cost (mg/L as N) 2,10 1,410 cost (mg/L as N	ю	2,230	4	232	223	4
(2) 7.3 7.0 4 12.9 cot to 4.5, lab (mg/L-as CaCO ₃) 314 314 0 640 26.0 25.1 4 380 26.0 25.1 4 380 28 28 0 8 15.6 15.0 4 21.3 180°C, dissolved (mg/L) 1,590 1,530 1 20,300 cnts (mg/L) 1,440 1,410 2 18,600 notal (mg/L-as N) <0.0	_	2,000	'n	286	172	vi
cot to 4.5, lab (mg/L.as CaCO ₃) 314 314 0 640 770 762 1 12,900 260 25.1 4 380 28 28 0 4.32 166 16.0 4 21.3 180°C, dissolved (mg/L) 1,550 1,530 1,410 20,300 cnts (mg/L) 1,440 1,410 2 18,600 soft ms/L as N) 2029 20 13.1 dissolved (mg/L.as N) 217 1,61 dissolved (mg/L.as N) 217 1,61 cot 1,630 cot 1	4	12.6	М	7.3	16.0	8
770 762 1 12,900 260 25.1 4 380 4 4 6 88 28 28 0 4.32 166 16.0 4 21.3 180°C, dissolved (mg/L) 1,590 1,530 1,410 20,300 cmts (mg/L) 1,440 1,410 2 18,600 smg/Las N) <0.029 <0.029 dissolved (mg/Las N) .17 .16 7 1,41 dissolved (mg/Las N) <10 <10 E.04	0	629	0	496	486	61
25.0 25.1 4 380 4 4 4 0 8 28 28 0 4.32 15.6 16.0 4 21.3 18.0°C, dissolved (mg/L, 1,550 1,530 1,410 20,300 cmts (mg/L, 1,110 20,300 cmts (mg/L,	-	13,100	çı	2,520	2,530	0
L) 1550 1530 0 4.32 L) 1550 1530 1 20,300 L) 1,440 1,410 2 18,600 3.80 3.80 0 13.1						

[RPD, relative percent difference, which is defined as [(sample value-replicate value+ replicate value+ replicate value) [2] × 100; relative percent difference was calculated on unrounded data; --, not analyzed; ND, not determined because data were less than the minimum reporting limit or estimated; <, less than; E, value estimated by laboratory] Table 15. Comparison of water-quality data for replicate and regular ground-water samples collected near Deer Trail, Colorado, 2000—Continued

	Well number		DIX3			90			025	
	Date		04/11/00			04/13/00			00/90/20	
	Time	1015	1016		0945	0960		1500	1505	
Property or constituent		Sample	Replicate	RPD	Sample	Replicate	RPD	Sample	Replicate	RPD
Aluminum, dissolved (Ilg/L as Al)		9>	>	ND	1	1	0	-	-	0
Antimony, dissolved (µg/L as Sb)		⊽	⊽	0	7	⊽	0	7	7	0
Arsenic, dissolved (µg/L as As)		å	۵	0	¹ Ε9	187	Q	en	m	0
Barium, dissolved (µg/L as Ba)		23	13	0	9	9	0	17	17	0
Beryllium, dissolved (µg/L as Be)		⊽	7	0	7	⊽	0	7	7	0
Botton, dissolved (Itg/L.ns B)		220	220	0	E750	750	Q.	440	410	7
Cadmium, dissolved (µg/L as Cd)		<1.0	<1.0	0	<1.0	0.10	0	<1.0	<1.0	0
Chromium, dissolved (µg/L ns Cr)		8	⊽	Q	N.	0.15	Q	N.	E.6	S
Cobalt, dissolved (µg/L as Co)		⊽	~	0	9	9	0	6	e	0
Copper, dissolved (µg/L as Cu)		М	М	0	61	63	ņ	10	6	Ξ
Iron, dissolved (µg/L as Fe)		OI>	<10 <10	0	≪200	00€>	0	€30	<30	0
Lead, dissolved (µg/L as Pb)		⊽	⊽	0	⊽	⊽	0	7	7	0
Manganese, dissolved (µg/L as Mn)		⊽	7	0	3,680	3,770	51	3,290	2,880	13
Mercury, dissolved (µg/L as Hg)		7	\$	0	۵	4	0	<.2	<2	0
Molybdenum, dissolved (µg/L as Mo)		⊽	7	0	m	3	0	6	6	0
Nickel, dissolved (µg/L as Ni)		9	9	0	6	6	0	33	14	8.1
Selenium, dissolved (µg/L as Se)		4	13	vo.	E17	ES	ON.	57	3	-18
Silver, dissolved (µg/L as Ag)		⊽	⊽	0	⊽	⊽	0	7	7	0
Strontium, dissolved (Jlg/L as Sr)		2,770	2,800	7	17,500	16,900	m	3,370	3,230	4
Zinc, dissolved (µg/L as Zn)		۵	⊽	R	63	99	s.	'n	9	-18
Uranium, natural (Ag/L as U)		28	28	0	164	164	0	7	4	0
Plutonium-238, whose water (pGAL as Pu)	(n	1	-1	1	i i		1		1	1
1 1 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0										

[RPD, relative percent difference, which is defined as [(sample value-replicate value)/((sample value + replicate value)/2)] × 100, relative percent difference was calculated on uncounded data; --, not analyzed; ND, not determined because data were less than the minimum reporting limit or estimated; <, less than; E, value estimated by laboratory] Comparison of water-quality data for replicate and regular ground-water samples collected near Deer Trail, Colorado, 2000—Continued Table 15.

						670			8	
	Date		02/11/00			10/10/00			10/12/00	
	Time	1345	1350		1445	1500		1300	1306	
Property or constituent	0.575550	Sample	Replicate	RPD	Sample	Replicate	RPD	Sample	Replicate	RPD
Specific conductance, lab (µS/cm at 25 °C)	G	16,000	16,000	0	4,900	4,910	0	16,200	16,100	-
pH, WH, laboratory (standard units)		7.2	7.2	0	7.2	7.2	0	7.1	7.1	0
Hardness total (mg/L as CaCO ₃)		10,000	10,000	0	2,900	2,900	0	10,000	10,000	0
Calcium, dissolved (mg/L as Ca)		414	417	7	794	7.67	0	437	439	0
Magnesium, dissolved (mg/L as Mg)		2,170	2,180	0	219	220	0	2,190	2,200	0
Sodium, dissolved (mg/L as Na)		2,070	2,070	0	334	334	0	2,110	2,100	0
Potassium, dissolved (mg/L as K)		12.0	12.7	'n	50	8.2	-	10.4	10.0	4
Acid neutralizing capacity, titration to 4.5, lab (mg/L as	, lab (mg/L as $\mathrm{CaCO}_3)$	638	629	0	385	473	21	640	640	0
Sulfate, dissolved (mg/L as SO ₄)		13,100	13,200	7	2,860	2,850	0	13,100	13,300	Ç
Chloride, dissolved (mg/L as Cl)		404	403	0	81.5	79.1	9	419	418	0
Fluoride, dissolved (mg/L as F)		οQ	Ľ.	6	1.0	1.0	0	ρď	ωq	0
Bromide, dissolved (mg/L as Br)		3.95	3.94	0	1.38	1.39	7	4.15	4.14	0
Silica, dissolved (mg/L as SiO ₂)		20.6	20.9	7	28.8	29.0	7	19.8	19.6	-
Solids, residue on evaporation at 180°C, dissolved (mg/L)	lissolved (mg/L)	21,100	21,200	0	5,270	5,170	7	21,500	20,900	m
Dissolved solids, sum of constituents (mg/L)	/L)	18,700	18,700	0	4,710	4,620	73	18,700	19,000	çı
Nitrite plus nitrate (mg/L.ns N)		14.4	15.5	-7	4.9	5.0	5	14.6	14.6	0
Nitrogen ammonia, dissolved (mg/L as N)	ê	<.030	<.030	0	110	Ξ.	0	.131	.135	φ
Nitrogen ammonia plus organic, total (mg/L as N)	g/Las N)	1.4	1.4	0	88	.73	19	1.5	1.5	0
Nitrogen ammonia plus organic, dissolved (mg/L as N)	d (mg/Las N)	1.5	1.5	0	Π,	83	-153	17	86	12
Phosphorus, total (mg/L as P)		E04	<00>	Q.	.12	Π.	v	EOH	<.05	$\frac{\Omega}{N}$
Phosphorus, dissolved (mg/L as P)		E03	E04	Q.	Π.	Ε.	0	<.05	E.04	R
Aluminum dissulated (Red as Al)		•								

[RPD, relative percent difference, which is defined as [(sample value-replicate value)/((sample value + replicate value)/2)] × 100, relative percent difference was calculated on uncounded data; --, not arralyzed; ND, not determined because data were less than the minimum reporting limit or estimated; <, less than; E, value estimated by laboratory] Table 15. Comparison of water-quality data for replicate and regular ground-water samples collected near Deer Trail, Colorado, 2000—Continued

	Wellnumber		8			D25			90 D	
	Date		02/11/00			10/10/00			10/12/00	
	Time	1345	1350		1445	1500		1300	1305	
Property or constituent		Sample	Replicate	RPD	Sample	Replicate	RPD	Sample	Replicate	RPD
Antimony, dissolved (Ug/L as Sb)		₽	D	0	۵	4	0	~	⊽	0
Arsenic, dissolved (Ilg/L as As)		Ξ	Ξ	0	e	8	0	6	6	0
Batium, dissolved (µg/L as Ba)		0	0	0	23	33	0	'n	w	0
Beryllium, dissolved (µg/L as Be)		0	0	0	۵	۵	0	7	⊽	0
Boron, dissolved (Jlg/L as B)		820	800	77	400	430	φ	970	970	0
Cadmium, dissolved (µg/L as Cd)		0.0	0.70	0	2.0	2.0	0	Т	т.	0
Chromium, dissolved (µg/L as Cr)		× ×	×.8	0	4.4	4.6	4	× ×	ж. У	0
Cobalt, dissolved (Itg/L.as Co)		0	∇	0	50	8	0	9	7	-15
Copper, dissolved (Jlg/L as Cu)		3.5	28	22	6	6	0	33	35	9-
Iron, dissolved (µg/L as Fe)		<200	<200	0	30	99	0	8	<150	0
Lead, dissolved (Itg/L as Pb)		∇	0	0	۵	۵	0	7	EOH	N
Manganese, dissolved (µg/L as Mn)		3,970	3,450	14	3,790	3,870	çı	3,760	3,730	-
Mercury, dissolved (µg/L as Hg)		E.2	<2	Ñ	<.2	\$	0	2	\$	0
Molybdenum, dissolved (µg/L as Mo)		5	0	0	14	13	7	4	ъ	29
Nickel, dissolved (Ug/L as Ni)		20	21	ņ	14	12	15	м	Ξ	-33
Selenium, dissolved (Itg/L as Se)		6	00	00	9	9	0	15	4	7
Silver, dissolved (Itg/L as Ag)		0	∇	0	۵	۵	0	7	⊽	0
Strontium, dissolved (µg/L as Sr)		16,500	16,600	7	3,550	3,550	0	16,500	16,500	0
Zinc, dissolved (µg/L as Zn)		42	49	-15	9	9	0	22	24	6-
Uranium, natural (µg/L.as U)		151	146	E)	8	28	6	95	171	$\overline{}$
Plutonium-238, whole water (pCi/L as Pu)	_	1	1	1	0	0.002	-200	0	0	0
Plutonium-239-240, whole water (pCi/L as Pu)	is Pu)	1	1	1	0.003	-0.002	1,000	1000	-0.002	999

Table 16. Statistical comparison of median concentrations for selected chemical constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000, and lowest applicable water-quality standard

Well	Sample size for statistical comparison	Minimum ground-water value used for statistical comparison ¹	Maximum ground-water value used for statistical comparison ¹	Median ground-water value used for statistical comparison ¹	Colorado standard	Type of regulatory standard	Probability that the median concentration exceeded the regulatory standard ²
			Nitrate ³ (rr	ig/L as N)			
D6	8	11	14.6	12	10	H	0.9961
D13	7	.018	.049	.025	10	H	.0078
D17	8	.679	5.68	2.60	10	H	.0039
D25	7	2.2	7.4	3.4	10	H	.0078
D29	8	.018	.03	.023	10	H	.0039
D30	8	.018	.06	.018	10	H	.0039
DTX1	7	1.1	2.8	1.29	10	H	.0078
DTX10A	8	.018	.025	.018	10	H	.0039
DTX2	8	.018	.05	.022	10	H	.0039
DTX3	8	.018	4.3	3.3	10	H	.0039
DTX4	8	.018	6.52	.241	10	H	.0039
DTX5	8	.018	.1	.018	10	H	.0039
DTX6	8	.22	.32	.25	10	H	.0039
DTX8A	8	.018	.06	.018	10	H	.0039
			Arsenic ⁴ (μ	g/L as As)			
D6	8	1	3.2	2.2	5	H	.0039
D13	8	.0.5	1	1	5	H	.0039
D17	8	1	2	1.35	5	H	.0039
D25	8	2	6	2.7	5	H	.0352
D29	8	.05	2	1	5	H	.0039
D30	8	.05	2.7	1	5	H	.0039
DTX1	8	1	3.5	1.95	5	H	.0039
DTX10A	8	.05	1	1	5	H	.0039
DTX2	8	.05	2	1.15	5	H	.0039
DTX3	8	.05	1	1	5	H	.0039
DTX4	8	.05	2	1	5	Н	.0039
DTX5	8	.Ω5	1	1	5	Н	.0039
DTX6	8	.05	2.2	1	5	H	.0039
DTX8A	8	.05	1	1	5	Н	.0039
			Cudmium ⁵ ()	ng/L as Cd)			
D6	8	.1	3.5	2	5	H	.0039
D13	8	0.022	0.5	0.5	5	H	.0039
D17	8	.05	.5	.5	5	Н	.0039

Table 16. Statistical comparison of median concentrations for selected chemical constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000, and lowest applicable water-quality standard—Continued

Well	Sample size for statistical comparison	Minimum ground-water value used for statistical comparison ¹	Maximum ground-water value used for statistical comparison ¹	Median ground-water value used for statistical comparison	Colorado standard	Type of regulatory standard	Probability that the median concentration exceeded the regulatory standard ²
		-	Cadmium ⁵ (µg/L a	s Cd)Continued			
D25	8	.5	1.5	1	5	Н	.0039
D29	8	.038	1	1	5	H	.0039
D30	8	.037	1	.5	5	Н	.0039
DTX1	8	.5	1	1	5	H	.0039
DTX10A	8	.05	1	.75	5	H	.0039
DTX2	8	.5	1	1	5	H	.0039
DTX3	8	.05	.5	.5	5	Н	.0039
DTX4	8	.1	1	1	5	Н	.0039
DTX5	8	.05	1	1	5	Н	.0039
DTX6	8	.1	1	1	5	Н	.0039
DTX8A	8	.05	.5	.5	5	Н	.0039
			Chromium (μg/L as Cr)			
D6	8	.4	18	.5	100	H, A	.0039
D13	8	.4	11	.45	100	H, A	.0039
D17	8	.4	3	.4	100	H, A	.0039
D25	8	.4	27	1.55	100	H, A	.0039
D29	8	.4	18	.5	100	H, A	.0039
D30	8	.4	9.5	.95	100	H, A	.0039
DTX1	8	.4	14	.5	100	H, A	.0039
DTX10A	8	.4	5.7	.75	100	H, A	.0039
DTX2	8	.4	11	.7	100	H, A	.0039
DTX3	8	.4	13	.5	100	H, A	.0039
DTX4	8	.4	11	.75	100	H, A	.0039
DTX5	8	.4	8.2	95	100	H, A	.0039
DTX6	8	.4	11	1.1	100	H, A	.0039
DTX8A	8	.4	.8	.45	100	H, A	.0039

Table 16. Statistical comparison of median concentrations for selected chemical constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000, and lowest applicable water-quality standard—Continued

Well	Sample size for statistical comparison	Minimum ground-water value used for statistical comparison ¹	Maximum ground-water value used for statistical comparison ¹	Median ground-water value used for statistical comparison ¹	Colorado standard	Type of regulatory standard	Probability that the median concentration exceeded the regulatory standard ²
			Copper (με	/L us Cu)			
D6	8	5	61	31	200	A	.0039
D13	8	2	4	2.5	200	A	.0039
D17	8	.05	1	.275	200	A	.0039
D25	8	7	11	9	200	A	.0039
D29	8	6	10	7.5	200	Α	.0039
D30	8	5	18	8.5	200	A	.0039
DTX1	8	6	14	8.5	200	A	.0039
DTX10A	8	3	6	5.5	200	A	.0039
DTX2	8	5	11	7	200	A	.0039
DTX3	8	2	5	3	200	A	.0039
DTX4	8	4	8	6	200	A	.0039
DTX5	8	4	8	6.5	200	A	.0039
DTX6	8	6	16	8.5	200	A	.0039
DTX8A	8	2	4	2	200	A	.0039
			Lend ⁶ (μg/	L as Pb)			
D6	8	.5	3.5	2	50	H	.0039
D13	8	.5	.5	.5	50	Н	.0039*
D17	8	.5	.5	.5	50	H	.0039*
D25	8	.5	1.5	1	50	H	.0039
D29	8	.5	1	1	50	H	.0039
D30	8	.5	1	.5	50	H	.0039
DTX1	8	.5	1	1	50	Н	.0039
DTX10A	8	.5	1	.75	50	Н	.0039
DTX2	8	.5	1	1	50	Н	.0039
DTX3	8	.5	.5	.5	50	Н	.0039*
DTX4	8	.5	1	1	50	Н	.0039
DTX5	8	.5	1	1	50	Н	.0039
DTX6	8	.5	1	1	50	H	.0039
DTX8A	8	.5	.5	.5	50	Н	.0039*

Table 16. Statistical comparison of median concentrations for selected chemical constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000, and lowest applicable water-quality standard—Continued

Well	Sample size for statistical comparison	Minimum ground-water value used for statistical comparison ¹	Maximum ground-water value used for statistical comparison	Median ground-water value used for statistical comparison ¹	Colorado standard	Type of regulatory standard	Probability that the median concentration exceeded the regulatory standard ²
			Mercury (µ	g/L us Hg)			
D6	8	.05	.2	.1	2	Н	.0039
D13	8	.05	.1	.1	2	Н	.0039
D17	8	.05	.1	.1	2	Н	.0039
D25	8	.05	.1	.1	2	Н	.0039
D29	8	.05	.1	.1	2	Н	.0039
D30	8	.05	.1	.1	2	Н	.0039
DTX1	8	.05	.1	.1	2	Н	.0039
DTX10A	8	.05	.1	.1	2	Н	.0039
DTX2	8	.05	.1	.1	2	Н	.0039
DTX3	8	0.05	0.1	0.1	2	Н	.0039
DTX4	8	.05	.1	.1	2	Н	.0039
DTX5	8	.05	.1	.1	2	Н	.0039
DTX6	8	.05	.1	.1	2	Н	.0039
DTX8A	8	.05	.1	.1	2	Н	.0039
			Molybde mum ⁷	(µg/L as Mo)			
D6	8	.5	4	3.25	6		
D13	8	.5	1	1	6		
D17	8	6	7	6	6		
D25	8	9	14	10.5	6		
D29	8	.5	1	1	6		
D30	8	1	4	2.5	6		
DTX1	8	5	7	5.5	6		
DTX10A	8	.5	3	1	6		
DTX2	8	1	2	1	6		
OTX3	8	.5	1	1	6		
DTX4	8	1	1	1	6		
DTX5	8	1	1	1	6		
DTX6	8	.5	1	1	6		
DTX8A	8	.5	2	.5	6		

Table 16. Statistical comparison of median concentrations for selected chemical constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000, and lowest applicable water-quality standard—Continued

Well	Sample size for statistical comparison	Minimum ground-water value used for statistical comparison	Maximum ground-water value used for statistical comparison ¹	Median ground-water value used for statistical comparison ¹	Colorado standard	Type of regulatory standard	Probability that the median concentration exceeded the regulatory standard ²
			Nickel (µg	/L us Ni)			
D6	8	6	23	12	100	Н	.0039
D13	8	2	10	4	100	Н	.0039
D17	8	.5	2	1	100	Н	.0039
D25	8	4	38	16.5	100	H	.0039
D29	8	.5	24	15	100	H	.0039
D30	8	2	15	12	100	H	.0039
DTX1	8	4	31	16	100	H	.0039
DTX10A	8	1	21	6.5	100	Н	.0039
DTX2	8	2	26	14	100	Н	.0039
DTX3	8	2	9	3.5	100	H	.0039
DTX4	8	.5	20	12	100	H	.0039
DTX5	8	6	46	11.5	100	Н	.0039
DTX6	8	.5	40	6	100	H	.0039
DTX8A	8	0.5	6	3	100	Н	.0039
			Selenium (p	g/L us Se)			
D6	8	6	17	8.4	20	A	.0039
D13	8	.5	1.7	1.1	20	A	.0039
D17	8	7.5	9	8.3	20	A	.0039
D25	8	1	6	2.7	20	A	.0039
D29	8	.5	3.3	1.35	20	A	.0039
D30	8	.5	4.9	1.2	20	A	.0039
DTX1	8	1	5.9	2.55	20	A	.0039
DTX10A	8	.5	2	1.2	20	A	.0039
DTX2	8	.5	4.3	1.55	20	A	.0039
DTX3	8	4	16.5	14.05	20	A	.0039
DTX4	8	1	11.6	2	20	Α	.0039
DTX5	8	.5	3.3	1.2	20	Α	.0039
DTX6	8	.5	6.1	3.35	20	Α	.0039
DTX8A	8	.5	1.2	1.1	20	Α	.0039

Table 16. Statistical comparison of median concentrations for selected chemical constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000, and lowest applicable water-quality standard—Continued

Well	Sample size for statistical comparison	Minimum ground-water value used for statistical comparison ¹	Maximum ground-water value used for statistical comparison	Median ground-water value used for statistical comparison	Colorado standard	Type of regulatory standard	Probability that the median concentration exceeded the regulatory standard ²
3 9			Zinc (µg/	L us Zn)			
D6	8	5	63	30	2,000	A	.0039
D13	8	.5	3	1.75	2,000	A	.0039
D17	8	.5	.5	.5	2,000	A	.0039*
D25	8	5	25	7	2,000	A	.0039
D29	8	8	19	11	2,000	A	.0039
D30	8	4	10	6.5	2,000	A	.0039
DTX1	8	4	8	6	2,000	A	.0039
DTX10A	8	2	10	4.5	2,000	A	.0039
DTX2	8	4	9	6	2,000	A	.0039
DTX3	8	1	2.5	2	2,000	A	.0039
DTX4	8	3	6	4	2,000	A	.0039
DTX5	8	3	7	4	2,000	A	.0039
DTX6	8	5	7	6	2,000	A	.0039
DTX8A	8	1.5	2	2	2,000	A	.0039

¹Many of the values used in the statistical comparison were derived from concentrations that are less than the minimum reporting limit and therefore were set equal to one-half the minimum reporting limit.

²Value is 1 minus the p-value resulting from a one-tailed Sign Test (Helsel and Hirsch, 1995), which is used to indicate the level of statistical evidence that selected constituent concentrations are significantly greater than regulatory standards. A value close to 1.0 indicates more evidence that the median concentration exceeded the standard, whereas a value close to 0 indicates little evidence that the median concentration exceeded the standard. The percent confidence of the test can be determined by subtracting the p-value from 1 and multiplying by 100. For example, if the p-value is 0.100, 1-p is 0.900, so the median concentration is greater than the regulatory standard with 90-percent confidence.

Data compared to standard are for nitrite plus nitrate. Results indicate nitrite is a minor component.

During 2000, standard was a proposed maximum contaminant level.

⁵All data were less than laboratory minimum reporting limit, except for one value (D6). Minimum reporting limits were sometimes greater than the water-quality standard.

⁶All data were less than laboratory minimum reporting limit. Laboratory minimum reporting limit is less than the water-quality standard.

⁷No regulatory standard for this constituent.

Table 17. Statistical evaluation of monotonic time-series trend using the Kendall's tau correlation coefficient for selected constituents in ground-water samples collected near Deer Trail, Colorado, 1999-2000

[Only values in bold indicate statistically significant trends. For this report, the Kendall's tau statistic (Helsel and Hirsch, 1995) is used as an indicator of monotonic correlation between concentration and time. By this method, positive values of Kendall's tau indicate upward trends and negative values indicate downward trends. Kendall's tau is a number between -1 and 1 that indicates increasing strength of the correlation. A p-value is listed to indicate the level of significance of the correlation; --, not computed]

	Nit	rate	Ars	enic	Cadr	nium	Chron	nium	Cop	pper	L	ead
Well	Tau	p-value	Tau	p-value	Tau	p-value	Tau	p-value	Tau	p-value	Tau	p- value
D6	.786	.009	.107	.804	428	.174	643	.035	.214	.536	1	1
D13	.238	.548	.428	.174	1	1	571	.063	036	1.000	1	1
D17	.214	.536	428	.174	1	1	607	.048	.714	.019	1	1
D25	428	.230	071	.902	1	1	250	.458	.250	.458	1	1
D29	1	1	.071	.902	1	1	286	.386	036	1.000	1	1
D30	607	.048	.500	.108	1	1	500	.108	.071	.902	1	1
DTX1	.048	1.000	.071	.902	1	1	571	.063	.428	.173	1	1
DTX IOA	1	1	.357	.266	1	1	036	1.000	178	.621	1	1
DTX2	393	.216	.142	.710	1	1	607	.048	.393	.216	1	1
DTX3	214	.536	1	1	1	1	214	.536	.178	.621	1	1
DTX4	.500	.108	.036	1.000	1	1	286	.386	<.001	1.000	1	1
DTX5	464	.138	.428	.174	1	1	750	.013	.071	.902	1	1
DTX6	250	.458	.464	.138	1	1	.071	.902	.214	.536	1	1
DTX8A	607	.048	1	1	1	1	1	1	.107	.804	1	1

	Mer	cury	Molyb	denum	Nic	kel	Sele	nium	Zi	ne	
Well	Tau	p-value	Tau	p-value	Tau	p-value	Tau	p-value	Tau	p-value	
D6	1	1	.214	.536	107	.804	.464	.138	.071	.902	
D13	1	1	.428	.174	.071	.902	1	1	071	.902	
D17	1	1	107	.804	<.001	1.000	286	.386	1	1	
D25	1	1	250	.458	178	.621	.036	1.000	250	.458	
D29	.357	.266	178	.621	464	.138	.464	.138	.107	.804	
D30	1	1	<.001	1.000	250	.458	.428	.174	500	.108	
DTX1	1	1	.678	.026	.214	.536	.214	.536	.286	.386	
DTX IOA	1	1	428	.174	464	.138	.250	.458	357	.266	
DTX2	1	1	.286	.386	.143	.710	.464	.138	.071	.902	
DTX3	1	1	321	.322	.214	.536	.250	.458	.178	.621	
DTX4	1	1	<.001	1.000	393	.216	.643	.035	143	.710	
DTX5	1	1	<.001	1.000	143	.710	.428	.174	036	1.000	
DTX6	1	1	286	.386	321	.322	.250	.458	25	.458	
DTX8A	1	1	250	.458	178	.621	1	1	107	.804	

No concentrations were greater than the laboratory minimum reporting limit.

Table 18. Methods used to analyze streambed-sediment samples collected near Deer Trail, Colorado, 2000

[Samples were analyzed at the National Water Quality Laboratory in Derner, except for constituents footnoted; MRL, minimum reporting level; MDC, minimum detectable concentration (radiochemical samples); KP, inductively coupled plasma; GFAA, graphite furnace atomic absorption; AA, atomic absorption; ASF, automated segmented-flow spectrophotometry; *, not applicable; mg/kg, milligrams per kilogram; µg/g, micrograms per gram; g/kg, grams per kilogram; pCi/g, picocuries per gram]

Property or Constituent	Units	Analytical method	MRL or MDC
·		Nutrients	
Nitrogen, ammonia plus organic ¹	mg/kg	Colorimetry, ASF, salicylate-hypochlorite	
Phosphorus, total as P	mg/kg	Colorimetry, ASF, phosphomolybdate	40
		Trace elements	
Aluminum	µg/g	DCP	2.8
Atsenic	µg/g	GFAA	1
Cadmium	µg/g	AA	.1
Chromium	µg/g	AA	.4
Copper	µg/g	AA	2
Lead	µg/g	AA	.1
Mercury	μg/g	AA manual cold vapor	.01
Molybdenum	µg/g	AA	.1
Nickel	µg/g	AA	.1
Selenium	µg/g	AA, Hydride generation, ASF	1
Zinc	µg/g	AA	3.1
		Carbons	
Inorganic carbon	g/kg	Coulimetric	.2
Organic carbon	g/kg	Wet exidation	.2
Total carbon	g/kg	Induction futnace (Leco)	.1
		Radioactivity	
Gross alpha ²	pCi/g	Thorium-230	
Gross beta ²	pCi/g	Cesium-137	
Plutonium-238 ²	pCi/g	Alpha spectrometry	
Plutonium-239+240 ²	pCi/g	Alpha spectrometry	

Analyses done by Severn Trent Laboratory in Arvada, Colorado, MRL calculated for each sample.

² Analyses done by Severn Trent Laboratory in Richland, Washington, MDC calculated for each sample.

(regkg, milligrams per kilogram; µg/g, micrograms per gram; g/kg, grams per kilogram; pCi/g, picocunies per gram; < less than; ±, plus or minus the analytical uncertainty] Table 19. Streambed-sediment data collected near Deer Trail, Colorado, July 17, 2000

ej. S	Date	Time	Total nitrogen (mg/kgj	Phos- phorus (mg/kg)	Aluminum (µg/g)	Arsenie (µg/g)	Cadmium (µg/g)	Chromium (µg/g)	Copper (µg/g)	Lead (µg/g)	Mercury (µg/g)	Molyb- denum (µg/g)
Biosolids-applied basin	00/11/00	1500	824	280	8,500	⊽	0.11	8.0	12.3	13.1	2070	0.18
Control basin (no biosolids)	02/11/20	1645	92	240	9.240	⊽	51.	10.2	10.5	13.8	8	8
Control basin, replicate	00/11/00	1705	730	009	10,400	⊽	14	10.8	12.8	15.4	8	15
Site	Date	Time	Nickel (µg/g)	Selenium (µg/g)	Zinc (µg/g)	Carbon, inorganic (g/kg)	Carbon, organic (g/kg)	Gross alpha (pCi/g)	Gross beta (pGi/g)	Plutoniu-238 (pCi/g)	Plutonium- 239+240 (pCi/g)	
Biosolids-applied 07/17/00 basin	00/11/00	1500	10.3	⊽	40.8	4.7	7.4	9.8 ± 4.4	44.3±7.7	0 ± 0.002	0.002 ± -0.003	
Control basin (no 07/17/00 biosolids)	00/11/00	1645	12.5	⊽	44.4	4	7.2	10.4 ± 4.7	44.2±7.8	0 ± 003	.002 ± .004	
Control basin, replicate	07/17/00	1705	12.6	⊽	48.0	\$	8.2	16.9 ± 5.9	44.1±7.6	.001 ± .002	.003 ± .003	